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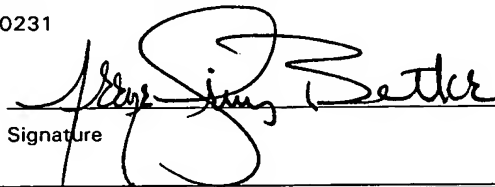
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U.S. PATENT APPLICATION

for

CALCILYTIC COMPOUNDS

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DESCRIPTION

CALCILYTIC COMPOUNDS

RELATED APPLICATIONS

5 This application is a divisional application of U.S. Serial No. 09/132,179,
now U.S. Patent No. 6,521,667
filed August 11, 1998, which is a continuation-in-part of U.S. Serial No.
08/629,608, filed April 9, 1996, now abandoned,
benefit of U.S. Provisional Application No. 60/032,263, filed December 3, 1996. which claims

FIELD OF THE INVENTION

The present invention relates to compounds able to inhibit calcium
receptor activity and the use of such compounds. Preferably, the compounds
described herein are administered to patients to achieve a therapeutic effect.

BACKGROUND OF THE INVENTION

Certain cells in the body respond not only to chemical signals, but also to
ions such as extracellular calcium ions (Ca^{2+}). Extracellular Ca^{2+} is under tight
homeostatic control and regulates various processes such as blood clotting, nerve
and muscle excitability, and proper bone formation.

Calcium receptor proteins enable certain specialized cells to respond to
changes in extracellular Ca^{2+} concentration. For example, extracellular Ca^{2+}
inhibits the secretion of parathyroid hormone (PTH) from parathyroid cells,
inhibits bone resorption by osteoclasts, and stimulates secretion of calcitonin from
C-cells.

PTH is the principal endocrine factor regulating Ca^{2+} homeostasis in the
blood and extracellular fluids. PTH, by acting on bone and kidney cells, increases
the level of Ca^{2+} in the blood. This increase in extracellular Ca^{2+} then acts as a
negative feedback signal, depressing PTH secretion. The

reciprocal relationship between extracellular Ca^{2+} and PTH secretion forms an important mechanism maintaining bodily Ca^{2+} homeostasis.

Extracellular Ca^{2+} acts directly on parathyroid cells to
 5 regulate PTH secretion. The existence of a parathyroid cell surface protein which detects changes in extracellular Ca^{2+} has been confirmed. (Brown et al., *Nature* 366:574, 1993.) In parathyroid cells, this protein, the calcium receptor, acts as a receptor for extracellular Ca^{2+} , detects changes in
 10 the ion concentration of extracellular Ca^{2+} , and initiates a functional cellular response, PTH secretion.

Extracellular Ca^{2+} can exert effects on different cell functions, reviewed in Nemeth et al., *Cell Calcium* 11:319, 1990. The role of extracellular Ca^{2+} in parafollicular (C-
 15 cells) and parathyroid cells is discussed in Nemeth, *Cell Calcium* 11:323, 1990. These cells were shown to express similar calcium receptors. (See, Brown et al., *Nature* 366:574, 1993; Mithal et al., *J. Bone Miner. Res.* 9, Suppl. 1, s282, 1994; Rogers et al., *J. Bone Miner. Res.* 9, Suppl. 1, s409, 1994; Garrett et al., *Endocrinology* 136:5202-5211, 1995.) The role of extracellular Ca^{2+} on bone osteoclasts is
 20 discussed by Zaidi, *Bioscience Reports* 10:493, 1990.

The ability of various molecules to mimic extracellular Ca^{2+} in vitro is discussed in references such as Nemeth et
 25 al., in "Calcium-Binding Proteins in Health and Disease," 1987, Academic Press, Inc., pp. 33-35; Brown et al., *Endocrinology* 128:3047, 1991; Chen et al., *J. Bone Miner. Res.* 5:581, 1990; and Zaidi et al., *Biochem. Biophys. Res. Commun.* 167:807, 1990.

30 Nemeth et al., PCT/US92/07175, International Publication Number WO 93/04373, Nemeth et al., PCT/US93/01642, International Publication Number WO 94/18959, and Nemeth et al., PCT/US94/12117, International Publication Number WO

95/11211, feature calcium receptor-active molecules and refer to calcilytics as compounds able to inhibit calcium receptor activity. For example, WO 94/18959 on page 8, lines 2-13 asserts:

5 Applicant is also the first to describe methods by which molecules active at these Ca^{2+} receptors can be identified and used as lead molecules in the discovery, development, design, modification and/or construction of useful calcimimetics or
10 calcilytics which are active at Ca^{2+} receptors. Such calcimimetics or calcilytics are useful in the treatment of various disease states characterized by abnormal levels of one or more components, e.g., polypeptides such as hormones,
15 enzymes or growth factors, the expresssion and/or secretion of which is regulated or affected by activity at one or more Ca^{2+} receptors.

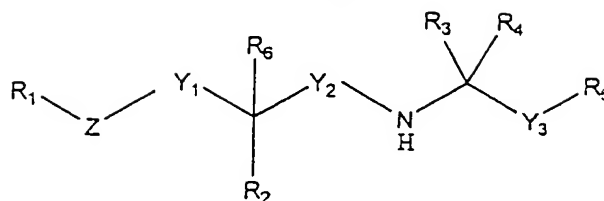
The references provided in the background are not admitted to be prior art to the pending claims.

20 SUMMARY OF THE INVENTION

The present invention features calcilytic compounds. "Calcilytic compounds" refer to compounds able to inhibit calcium receptor activity. The ability of a compound to "inhibit calcium receptor activity" means that the compound
25 causes a decrease in one or more calcium receptor activities evoked by extracellular Ca^{2+} .

The use of calcilytic compounds to inhibit calcium receptor activity and/or achieve a beneficial effect in a patient are described below. Also described below are
30 techniques which can be used to obtain additional calcilytic compounds.

An example of featured calcilytic compounds are Structure I α, α -disubstituted arylalkylamine derivatives having the chemical formula:

STRUCTURE I

where R_1 is selected from the group consisting of: aryl, longer-length alk, and cycloalk;

5 R_2 is selected from the group consisting of: lower alk, cycloalk, alkoxy, H, OH, =O, C(O)OH, C(O)O-lower alk, C(O)NH-lower alk, C(O)N(lower alk)₂, SH, S-lower alk, NH₂, NH-lower alk, and N(lower alk)₂;

10 R_3 and R_4 is each independently lower alk or together cyclopropyl;

R_5 is aryl;

R_6 if present is either hydrogen, lower alkyl or lower alkenyl, wherein R_6 is not present if R_2 is =O;

Y_1 is either covalent bond, alkylene, or alkenylene;

15 Y_2 is alkylene;

Y_3 is alkylene; and

Z is selected from the group consisting of: covalent bond, O, S, NH, N-lower alk, alkylene, alkenylene, and alkynylene, provided that if Z is either O, S, NH, or N-lower alk, then Y_1 is not a covalent bond, further provided that Y_1 and Z may together be a covalent bond;

and pharmaceutically acceptable salts and complexes thereof.

25 The terms aryl, longer-length alk, lower alk, cycloalk, alkoxy, alkylene, alkenylene, and alkynylene, along with possible substituents are defined in Section II, *infra*.

Section II, *infra*, also provides definitions for other chemical groups described in the present application.

Preferred calcilytic compounds have an $IC_{50} \leq 50 \mu M$, more preferably an $IC_{50} < 10 \mu M$, and even more preferably an $IC_{50} < 1 \mu M$, as measured using the "Calcium Receptor Inhibitor Assay" described in Example 1, *infra*.

Thus, a first aspect of the present invention features a method of treating a patient by administering to the patient a therapeutically effective amount of a Structure I α, α -disubstituted arylalkylamine derivative. Treatment can be carried out, for example, to retard the disease in a patient having a disease or to prophylactically retard or prevent the onset of a disease.

A therapeutically effective amount is the amount of compound which achieves a therapeutic effect by retarding a disease in a patient having a disease or prophylactically retarding or preventing the onset of a disease. Preferably, it is an amount which relieves to some extent one or more symptoms of a disease or disorder in a patient; returns to normal either partially or completely one or more physiological or biochemical parameters associated with or causative of the disease or disorder; and/or reduces the likelihood of the onset of the disease or disorder.

A "patient" refers to a mammal in which compounds characterized by their ability to inhibit calcium receptor activity, *in vivo* or *in vitro*, will have a beneficial effect. Preferably, the patient is a human being.

Patients benefiting from the administration of a therapeutic amount of a calcilytic compound can be identified using standard techniques known to those in the medical profession. Diseases or disorders which can be treated by inhibiting one or more calcium receptor activities include

one or more of the following types: (1) those characterized by an abnormal bone and mineral homeostasis; (2) those characterized by an abnormal amount of an extracellular or intracellular messenger whose production can be affected by one or more calcium receptor activities; (3) those characterized by an abnormal effect (e.g., a different effect in kind or magnitude) of an intracellular or extracellular messenger which can itself be ameliorated by one or more calcium receptor activities; and (4) other diseases or disorders where inhibition of one or more calcium receptor activities exerts a beneficial effect, for example, in diseases or disorders where the production of an intracellular or extracellular messenger stimulated by receptor activity compensates for an abnormal amount of a different messenger. Examples of extracellular messengers whose secretion and/or effect can be affected by inhibiting calcium receptor activity are believed to include inorganic ions, hormones, neurotransmitters, growth factors, and chemokines. Examples of intracellular messengers include cAMP, cGMP, IP₃, calcium, magnesium, and diacylglycerol.

Preferably, a patient is a human having a disease or disorder characterized by one or more of the following: (1) an abnormal bone or mineral homeostasis; (2) an abnormal amount of an extracellular or intracellular messenger which is ameliorated by a compound able to effect one or more calcium receptor activities; and (3) an abnormal effect of an intracellular or extracellular messenger which is ameliorated by a compound able to effect one or more calcium receptor activities.

Preferably, the disease or disorder is characterized by an abnormal bone and mineral homeostasis, more preferably calcium homeostasis. Abnormal calcium homeostasis is characterized by one or more of the following activities: (1)

an abnormal increase or decrease in serum calcium; (2) an abnormal increase or decrease in urinary excretion of calcium; (3) an abnormal increase or decrease in bone calcium levels, for example, as assessed by bone mineral density measurements; (4) an abnormal absorption of dietary calcium; (5) an abnormal increase or decrease in the production and/or release of messengers which affect serum calcium levels such as PTH and calcitonin; and (6) an abnormal change in the response elicited by messengers which affect serum calcium levels. The abnormal increase or decrease in these different aspects of calcium homeostasis is relative to that occurring in the general population and is generally associated with a disease or disorder.

Preferably, the calcilytic compounds are used to treat diseases or disorders selected from the group consisting of: hypoparathyroidism, osteosarcoma, periodontal disease, fracture healing, osteoarthritis, rheumatoid arthritis, Paget's disease, humoral hypercalcemia malignancy, and osteoporosis.

Another aspect of the present invention describes a method of treating a patient comprising the step of administering to the patient an amount of a calcilytic compound sufficient to increase serum PTH level. Preferably, the method is carried out by administering an amount of the compound effective to cause an increase in duration and/or quantity of serum PTH level sufficient to have a therapeutic effect.

Increasing serum PTH may be used to achieve a therapeutic effect by retarding a disease in a patient having the disease or prophylactically retarding or preventing the onset of a disease. Prophylactic treatment can be performed, for example, on a person with an abnormally low serum PTH; or on a person without a low serum PTH, but were increasing PTH

has a beneficial effect. An abnormally low serum PTH is a serum PTH level lower than that occurring in the general population, and is preferably an amount associated with a disease or the onset of a disease.

- 5 Increasing serum PTH levels can be used to treat different types of diseases including bone and mineral related diseases.

10 In different embodiments, the compound administered to a patient causes an increase in serum PTH having a duration up to one hour, about one to about twenty-four hours, about one to about twelve hours, about one to about six hours, about one to about five hours, about one to about four hours, about two to about five hours, about two to about four hours, or about three to about six hours.

- 15 In additional different embodiments, the compound administered to a patient causes an increase in serum PTH up to 0.5 fold, 0.5 to 5 fold, 5 fold to 10 ten fold, and at least 10 fold, greater than peak serum PTH in the patient. The peak serum level is measured with respect to the patient
20 not undergoing treatment.

Another aspect of the present invention features Structure I calcilytic compounds.

- Another aspect of the present invention features a pharmaceutical composition comprising a pharmaceutically
25 acceptable carrier and a calcilytic compound described herein. The pharmaceutical composition contains the calcilytic compound in a form suitable for administration into a mammal, preferably, a human being. Preferably, the pharmaceutical composition contains an amount of a calcilytic
30 compound in a proper pharmaceutical dosage form sufficient to exert a therapeutic effect on a human being. However, multiple doses of pharmaceutical compositions may be used to treat a patient.

Considerations and factors concerning dosage forms suitable for administration are known in the art and include potential toxic effects, solubility, route of administration, and maintaining activity. For example, pharmaceutical
5 compositions injected into the bloodstream should be soluble.

Another aspect of the present invention features a method of screening for Structure I α,α -disubstituted arylalkylamine derivatives able to inhibit calcium receptor activity. The method involves the steps of contacting a cell
10 having a calcium receptor with a Structure I α,α -disubstituted arylalkylamine derivative and measuring the ability of the compound to inhibit calcium receptor activity.

The screening method can be carried out *in vivo* or *in vitro* and is particularly useful to identify those Structure
15 I α,α -disubstituted arylalkylamine derivatives most able to act as calcilytic compounds. *In vivo* assays include measuring a physiological parameter related to calcium receptor activity, such as serum hormone levels or serum calcium ion concentration. *In vitro* assays include measuring
20 the ability of the calcilytic compound to affect intracellular calcium concentration, or cellular hormone secretion. Examples of hormones levels which can be affected by calcilytic compounds include PTH and calcitonin.

The calcilytic compounds described herein can be used as
25 part of *in vivo* or *in vitro* methods. Preferably, the compounds are used *in vivo* to achieve a beneficial effect in a patient. Examples of *in vitro* uses, and other *in vivo* uses, include use in a method to identify other calcilytic compounds and use as a tool to investigate calcium receptor
30 activity or the physiological effects of inhibiting calcium receptor activity in different organisms.

Other features and advantages of the invention will be apparent from the following detailed description of the invention, examples, and the claims.

DETAILED DESCRIPTION OF THE INVENTION

5 The present application demonstrates the ability of calcilytic compounds to exert a physiologically relevant effect on a cell by illustrating the ability of such compounds to increase PTH secretion and also identifies a target site for calcilytic compounds. The present
10 application is believed to be the first to demonstrate that calcilytic compounds can increase PTH secretion.

Calcium receptors are present on different cell types and can regulate different responses in different cell types. While the calcilytic compounds described herein are believed
15 to act at a calcium receptor through a calcium receptor-activity modulating site, unless otherwise explicitly stated in the claims that a compound exerts an effect by acting at a calcium receptor through such a site, there is no intention to limit the claimed methods or compound to requiring
20 inhibition of calcium receptor activity or any particular mode of action. Rather, the present application demonstrates that compounds able to inhibit calcium receptor activity, whose calcilytic activity can be measured *in vivo* or *in vitro*, exert significant physiological effects. For example,
25 the present application demonstrates the ability of different calcilytic compounds to prevent Ca^{2+} inhibition of PTH and, thereby, result in an increase in PTH release.

Compounds binding at the calcium receptor-activity modulating site can be identified using a labeled compound
30 binding to the site in a competition-binding assay format.

Preferred calcilytic compounds described herein are Structure I α, α -disubstituted arylalkylamine derivatives able

to inhibit calcium receptor activity. Other aspects of the present invention include assays which can be used to identify those Structure I α,α -disubstituted arylalkylamine derivatives expected to be effective in inhibiting calcium
 5 receptor activity, and/or exerting a therapeutic effect in a patient; preferred groups of Structure I α,α -disubstituted arylalkylamine derivatives; and the use of the compounds described herein to treat different diseases or disorders..

I. Calcium Receptor Activity

10 Calcium receptors respond to changes in extracellular calcium levels. The exact changes resulting from calcium receptor activity depend on the particular receptor and the cell containing the receptor. For example, the *in vitro* effect of calcium on the calcium receptor in a parathyroid
 15 cell includes the following:

1. An increase in internal calcium $[Ca^{2+}]_i$. The increase is due to the influx of external calcium and/or to the mobilization of internal calcium. Characteristics of the increase in internal calcium include the following:
 - 20 (a) A rapid (time to peak < 5 seconds) and transient increase in $[Ca^{2+}]_i$ that is refractory to inhibition by $1 \mu M$ La^{3+} or $1 \mu M$ Gd^{3+} and is abolished by pretreatment with ionomycin (in the absence of extracellular Ca^{2+});
 - (b) The increase is not inhibited by di-
 25 hydropyridines;
 - (c) The transient increase is abolished by pretreatment for 10 minutes with 10 mM sodium fluoride;
 - (d) The transient increase is diminished by pretreatment with an activator of protein kinase C (PKC),
 30 such as phorbol myristate acetate (PMA), mezerein or (-)-indolactam V. The overall effect of the protein kinase C activator is to shift the concentration-response curve of

calcium to the right without affecting the maximal response;
and

(e) Pretreatment with pertussis toxin (100 ng/ml for > 4 hours) does not affect the increase.

5 2. A rapid (< 30 seconds) increase in the formation of inositol-1,4,5-triphosphate and/or diacylglycerol. Pretreatment with pertussis toxin (100 ng/ml for > 4 hours) does not affect this increase;

10 3. The inhibition of dopamine- and isoproterenol-stimulated cyclic AMP formation. This effect is blocked by pretreatment with pertussis toxin (100 ng/ml for > 4 hours); and

15 4. The inhibition of PTH secretion. Pretreatment with pertussis toxin (100 ng/ml for > 4 hours) does not affect the inhibition of PTH secretion.

Calcilytic activity of a compound can be determined using techniques such as those described in the examples below and those described in publications such as Nemeth et al., PCT/US92/07175, International Publication Number WO 93/04373, Nemeth et al., PCT/US93/01642, International Publication Number WO 94/18959, and Nemeth et al., PCT/US94/12117, International Publication Number WO 95/11211 (each of which are hereby incorporated by reference herein).

25 Calcilytic activity varies depending upon the cell type in which the activity is measured. For example, calcilytic compounds possess one or more, and preferably all, of the following characteristics when tested on parathyroid cells *in vitro*:

30 1. The compound blocks, either partially or completely, the ability of increased concentrations of extracellular Ca^{2+} to:

(a) increase $[Ca^{2+}]_i$,

(b) mobilize intracellular Ca^{2+} ,

(c) increase the formation of inositol-1,4,5-triphosphate,

(d) decrease dopamine- or isoproterenol-stimulated cyclic AMP formation, and

5 (e) inhibit PTH secretion;

2. The compound blocks increases in Cl^- current in *Xenopus* oocytes injected with poly(A)⁺-mRNA from bovine or human parathyroid cells elicited by extracellular Ca^{2+} , but not in *Xenopus* oocytes injected with water; and

10 3. Similarly, the compound blocks a response in *Xenopus* oocytes, injected with cloned nucleic acid expressing the calcium receptor, elicited by extracellular Ca^{2+} or a calcimimetic compound (i.e., a compound able to mimic the effect of extracellular Ca^{2+} , including compounds potentiating
15 the effect of extracellular Ca^{2+}).

Calcium receptors are present in different cells. The pharmacological effects of the following cells, in response to extracellular Ca^{2+} , is consistent with the presence of a calcium receptor: parathyroid cell, bone osteoclast,
20 juxtaglomerular kidney cell, proximal tubule kidney cell, distal tubule kidney cell, central nervous system cell, peripheral nervous system cell, cell of the thick ascending limb of Henle's loop and/or collecting duct, keratinocyte in the epidermis, parafollicular cell in the thyroid (C-cell),
25 intestinal cell, trophoblast in the placenta, platelet, vascular smooth muscle cell, cardiac atrial cell, gastrin-secreting cell, glucagon-secreting cell, kidney mesangial cell, mammary cell, endocrine and exocrine cells in the pancreas, fat/adipose cell, immune cell, GI tract cell, skin
30 cell, adrenal cell, pituitary cell, hypothalamic cell and cell of the subfornical organ.

The presence of a calcium receptor on the following cells have been confirmed using physical data, such as

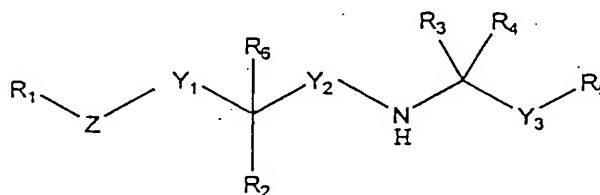
hybridization with nucleic acid encoding a calcium receptor:
parathyroid cell, central nervous system cell, peripheral
nervous system cell, cell of the thick ascending limb of
Henle's loop and/or collecting duct in the kidney,

- 5 parafollicular cell in the thyroid (C-cell), intestinal cell,
GI tract cell, pituitary cell, hypothalamic cell, cell of the
subfornical organ, and endocrine and exocrine cells in the
pancreas.

II. α,α -Disubstituted Arylalkylamine Derivatives

- 10 Structure I α,α -disubstituted arylalkylamine
derivatives have the following chemical formula:

STRUCTURE I



- where R_1 is selected from the group consisting of: aryl,
15 longer-length alk, and cycloalk. Preferably, R_1 is either
optionally substituted phenyl, optionally substituted
pyridyl, optionally substituted benzothiopyranyl, optionally
substituted carbazole, optionally substituted naphthyl,
optionally substituted tetrahydronaphthyl, optionally
20 substituted longer-length alkyl, optionally substituted
longer-length alkenyl or optionally substituted cycloalk.

- More preferably, R_1 is either an optionally substituted
phenyl; an optionally substituted naphthyl; an optionally
substituted pyridyl; an optionally substituted
25 benzothiopyranyl; an optionally substituted carbazole;

unsubstituted longer-length alkyl; unsubstituted longer-length alkenyl; or monosubstituted longer-length alkyl or alkenyl, where the monosubstituent is either an optionally substituted phenyl or an optionally substituted cycloalkyl
 5 provided that the optionally substituted phenyl or optionally substituted cycloalkyl can have one to four substituents each independently selected from the group consisting of: alkoxy, lower-haloalkyl, S-unsubstituted alkyl, lower-haloalkoxy, unsubstituted alkyl, unsubstituted alkenyl, halogen, SH, CN,
 10 NO₂, NH₂, and OH;

R₂ is selected from the group consisting of: lower alk, cycloalk, alkoxy, H, OH, =O, C(O)OH, C(O)O-lower alk, C(O)NH-lower alk, C(O)N(lower alk)₂, SH, S-lower alk, NH₂, NH-lower alk, and N(lower alk)₂. More preferably, R₂ is OH or alkoxy,
 15 even more preferably, R₂ is OH or methoxy;

R₃ and R₄ is each independently lower alk or together cyclopropyl. Preferably, R₃ and R₄ are each independently a lower alkyl, more preferably, R₃ and R₄ are each independently methyl or ethyl;

20 R₅ is aryl. Preferably, R₅ is either optionally substituted naphthyl or optionally substituted phenyl. More preferably, R₅ is substituted phenyl having a substituent in the *meta* or *para* position and optionally containing additional substituents;

25 R₆ if present is either hydrogen, lower alkyl or lower alkenyl, wherein R₆ is not present if R₂ is =O. Preferably R₆ is either hydrogen or lower alkyl, more preferably R₆ is hydrogen.

Y₁ is either covalent bond, alkylene, or alkenylene.
 30 Preferably, Y₁ is either covalent bond or lower alkylene. More preferably, Y₁ is methylene;

Y₂ is alkylene. Preferably, Y₂ is lower alkylene. More preferably, Y₂ is methylene;

Y_1 is alkylene. Preferably, Y_1 is lower alkylene. More preferably, Y_1 is methylene;

Z is selected from the group consisting of: covalent bond, O, S, NH, N-lower alk, alkylene, alkenylene, and alkynylene, provided that if Z is either O, S, NH, or N-lower alk, then Y_1 is not a covalent bond, further provided that Y_1 and Z may together be a covalent bond. Preferably, Z is selected from the group consisting of: covalent bond, O, S, NH, N-lower alk, and alkylene. More preferably, Z is either O, S, lower alkylene, even more preferably, Z is O;

and pharmaceutically acceptable salts and complexes thereof.

"Alk" refers to either alkyl, alkenyl, or alkynyl.

"Lower alk" refers to either lower alkyl, lower alkenyl, or lower alkynyl, preferably, lower alkyl.

"Alkenyl" refers to an optionally substituted hydrocarbon group containing at least one carbon-carbon double bond between the carbon atoms and containing 2-15 carbon atoms joined together. The alkenyl hydrocarbon group may be straight-chain or contain one or more branches. Branched- and straight-chain alkenyl preferably have 2 to 7 carbons, each of which may be optionally substituted. Alkenyl substituents are each independently selected from the group consisting of: lower alkyl, lower alkenyl, halogen, alkoxy, lower haloalkyl, lower haloalkoxy, methylene dioxy, unsubstituted aryl, unsubstituted cycloalkyl, OH, SH, CN, NO, NO₂, NH₂, CH=NNHC(O)NH₂, CH=NNHC(S)NH₂, CH₂O-lower alkyl, C(O)lower alkyl, C(O)NH₂, C(O)NH-lower alkyl, C(O)N(lower alkyl)₂, C(O)OH, C(O)O-lower alkyl, NH-lower alkyl, N(lower alkyl)₂, NHC(O)unsubstituted aryl, NHC(O)lower alkyl, N=N-unsubstituted aryl, NHC(O)NH₂, N(lower alkyl)C(O)lower alkyl, NHC(S)lower alkyl, N(lower alkyl)C(S)lower alkyl, NHS(O)lower alkyl, N(lower alkyl)S(O)lower alkyl, OC(O)lower alkyl,

OCH₂C(O)OH, OC(S)lower alkyl, S(O)lower alkyl, SC(O)lower alkyl, S-lower alkyl, S-lower haloalkyl, SO₂-lower alkyl, SO₂-lower haloalkyl, S(O)₂NH₂, S(O)₂NH-lower alkyl, and S(O)₂N(lower alkyl)₂. Preferably, no more than three
 5 substituents are present. Even more preferably, the alkenyl is a lower alkenyl, which is an unsubstituted branched- or straight-chain alkenyl having 2 to 4 carbons.

"Alkyl" refers to an optionally substituted hydrocarbon group joined by single carbon-carbon bonds and having 1-15
 10 carbon atoms joined together. The alkyl hydrocarbon group may be straight-chain or contain one or more branches. Branched- and straight-chain alkyl preferably have 1 to 7 carbons, each of which may be optionally substituted. Alkyl substituents are each independently selected from the
 15 substituents described above for alkenyl. Preferably, no more than three substituents are present. More preferably, the alkyl is a lower alkyl, which is an unsubstituted branched- or straight-chain alkyl 1 to 4 carbons in length.

"Alkynyl" refers to an optionally substituted
 20 hydrocarbon group containing at least one carbon-carbon triple bond between the carbon atoms and containing 2-15 carbon atoms joined together. The alkynyl hydrocarbon group may be straight-chain or contain one or more branches. Branched- and straight-chain alkynyl preferably have 2 to 7
 25 carbons, each of which may be optionally substituted. Alkynyl substituents are each independently selected from the substituents described above for alkenyl. Preferably, no more than three substituents are present. More preferably, the alkynyl is a lower alkynyl, which is an unsubstituted
 30 branched- or straight-chain alkynyl having 2 to 4 carbons.

"Alkenylene" refers to an optionally substituted hydrocarbon chain containing at least one carbon-carbon

double bond between the carbon atoms. The alkenylene chain has 2 to 6 carbons and is attached at two locations to other functional groups or structural moieties. The alkenylene substituents are each independently selected from the substituents described above for alkenyl. Preferably, no more than three substituents are present. More preferably, the alkenylene is a "lower alkenylene," which is an unsubstituted branched- or straight-chain alkenylene having 2 to 3 carbons.

10 "Alkoxy" refers to oxygen joined to an unsubstituted alkyl 1 to 12 carbon atoms in length, preferably 1 to 2 carbons in length. More preferably, the alkoxy is methoxy.

"Alkylene" refers to an optionally substituted hydrocarbon chain containing only carbon-carbon single bonds between the carbon atoms. The alkylene chain has 1 to 6 carbons and is attached at two locations to other functional groups or structural moieties. The alkylene substituents are each independently selected from the substituents described above for alkenyl. Preferably, no more than three substituents are present. More preferably, the alkylene is a "lower alkylene," which is an unsubstituted branched-, or straight-chain alkylene having 1 to 3 carbons.

"Alkynylene" refers to an optionally substituted hydrocarbon chain containing at least one carbon-carbon triple bond between the carbon atoms. The alkynylene chain has 2 to 6 carbons and is attached at two locations to other functional groups or structural moieties. The alkynylene substituents are each independently selected from the substituents described above for alkenyl. More preferably, the alkynylene is a "lower alkynylene," which is an unsubstituted branched- or straight-chain alkynylene having 2 to 3 carbons.

"Aryl" refers to an optionally substituted aromatic group with at least one ring having a conjugated pi- electron system, containing up to two conjugated or fused ring systems. Aryl includes carbocyclic aryl, heterocyclic aryl and biaryl groups, all of which may be optionally substituted. Preferably, the aryl is either optionally substituted phenyl, optionally substituted pyridyl, optionally substituted benzothiopyranyl, optionally substituted carbazole, optionally substituted naphthyl, optionally substituted tetrahydronaphthyl.

Different substituents are preferred for the Structure I left hand R_1 aryl and the Structure I R_2 right hand aryl. Preferably, the aryl has no more than five independently selected substituents.

Preferably, when R_1 is an aryl, the aryl is either optionally substituted phenyl, optionally substituted pyridyl, optionally substituted benzothiopyranyl, optionally substituted carbazole, optionally substituted naphthyl, or optionally substituted tetrahydronaphthyl. Preferred, R_1 substituents are each independently selected from the group consisting of: unsubstituted alkyl, unsubstituted alkenyl, halogen, alkoxy, lower haloalkyl, lower haloalkoxy, methylene dioxy, unsubstituted aryl, unsubstituted cycloalkyl, OH, SH, CN, NO, NO_2 , NH_2 , methylene dioxy, CH=NNHC(O)NH_2 , CH=NNHC(S)NH_2 , CH_2O -unsubstituted alkyl, C(O) unsubstituted alkyl, C(O)NH_2 , C(O)NH -unsubstituted alkyl, C(O)N (unsubstituted alkyl) $_2$, C(O)OH , C(O)O -unsubstituted alkyl, NH -unsubstituted alkyl, N (unsubstituted alkyl) $_2$, NHC(O) unsubstituted aryl, NHC(O) unsubstituted alkyl, N=N -unsubstituted aryl, NHC(O)NH_2 , N (unsubstituted alkyl) C(O) unsubstituted alkyl, NHC(S) unsubstituted alkyl, N (unsubstituted alkyl) C(S) unsubstituted alkyl, NHS(O) unsubstituted alkyl, N (unsubstituted

alkyl)S(O)unsubstituted alkyl, NS(O)₂aryl, OC(O)unsubstituted alkyl, OCH₂C(O)OH, OC(S)unsubstituted alkyl, S(O)unsubstituted alkyl, SC(O)unsubstituted alkyl, S-unsubstituted alkyl, S-unsubstituted haloalkyl, SO₂-unsubstituted alkyl, SO₂-unsubstituted haloalkyl, S(O)₂NH₂, S(O)₂NH-unsubstituted alkyl, and S(O)₂N(unsubstituted alkyl)₂.

Preferred R₁ aryl substituents are each independently selected from the group consisting of: alkoxy, methylene dioxy, N(CH₃)₂, C(O)OCH₃, phenyl, lower-haloalkyl, S-unsubstituted alkyl, lower-haloalkoxy, unsubstituted alkyl, unsubstituted alkenyl, halogen, SH, CN, NO₂, NH₂, OH and sulfamoyl. More preferably, each R₁ aryl substituent is independently selected from the group consisting of: unsubstituted C₁-C₇ alkyl, C₁-C₇ alkoxy, lower haloalkoxy, CF₃, F, Cl, Br, CN, NO₂ and sulfamoyl.

In another preferred embodiment, R₁ is either 2-CN-phenyl, 2,3-dichloro phenyl, 2-nitro-phenyl, 2-cyano-3-chloro-phenyl, or 2,3-dichloro-4-sulfamoyl-phenyl.

R₂ right hand aryl substituents are each independently selected from the substituents described above for alkenyl. In a preferred embodiment, the R₂ aryl substituents are each independently selected from the group consisting of: methoxy, lower alkyl, lower haloalkoxy, CFH₂, CHF₂, CF₃, OCH₂CF₃, F, Cl, Br, I, OH, SH, CN, NO₂, NH₂, methylene dioxy, NH-lower alkyl, N(lower alkyl)₂, C(O)lower alkyl, S-lower alkyl, S(O)lower alkyl, S(O)₂lower alkyl, OC(O)lower alkyl, SC(O)lower alkyl, OC(S)lower alkyl, NHC(O)lower alkyl, N(lower alkyl)C(O)lower alkyl, NHC(S)lower alkyl, N(lower alkyl)C(S)lower alkyl, NHS(O)lower alkyl, N(lower alkyl)S(O)lower alkyl, C(O)OH, C(O)O-lower alkyl, C(O)NH₂, C(O)NH-lower alkyl, C(O)N(lower alkyl)₂, S(O)₂NH₂, S(O)₂NH-lower alkyl, and S(O)₂N(lower alkyl)₂.

In another preferred embodiment, R₃ aryl substituents are each independently selected from the group consisting of: methylene dioxy, methoxy, lower-haloalkyl, S-lower alkyl, lower-haloalkoxy, lower alkyl, halogen, SH, CN, OH, Cl, F, and Br. Preferred halogens are Cl, F, and Br.

"Carbocyclic aryl" refers to an aromatic ring or ring system having all carbon atoms. The carbon atoms are optionally substituted.

"Cycloalk" refers to an optionally substituted cyclic alkyl or an optionally substituted non-aromatic cyclic alkenyl and includes monocyclic and multiple ring structures such as bicyclic and tricyclic. The cycloalk has 3 to 15 carbon atoms, preferably, 5 to 12 carbon atoms. Optional substituents for the cycloalk are each independently selected from the group described above for alkenyl. Preferably, no more than three substituents are present. More preferably, the cycloalk is unsubstituted, even more preferably it is an unsubstituted cyclic alkyl. Preferred cycloalkyl groups include cyclohexyl and adamantyl.

"Haloalk" refers to substituted alkyl or substituted alkenyl, having no more than 4 carbons, where the substituents are halogens and at least one halogen is present. Preferably, the haloalk is an alkyl 1 to 3 carbons in length and the halogens are each independently either Cl or F, more preferably the alkyl has 2 carbons, more preferably the haloalkyl is a lower haloalkyl which has 1 carbon.

"Heterocyclic aryl" refers to an aryl having 1 to 3 heteroatoms as ring atoms in the aromatic ring and the remainder of the ring atoms are carbon atoms. Suitable heteroatoms include oxygen, sulfur, and nitrogen. Examples

of heterocyclic aryl include indolyl, pyridyl, quinolinyl, and isoquinolinyl.

"Longer-length alk" refers to either longer-length alkyl, longer-length alkenyl, or longer-length alkynyl; preferably, longer-length alkyl or longer-length alkenyl. More preferably a longer-length alk is 4 to 20 carbon atoms.

"Longer-length alkenyl" refers to an optionally substituted hydrocarbon group containing at least one carbon-carbon double bond between the carbon atoms, and which contains 2-20 carbon atoms joined together. Preferably, the longer-length alkenyl is 4 to 20 carbon atoms. The longer-length alkenyl hydrocarbon group may be straight-chain or contain one or more branches. Longer-length alkenyl substituents are each independently selected from the alkenyl substituent list described above. Preferably, the longer-length alkenyl is either unsubstituted or has one cycloalk or phenyl substituent. More preferably, the cycloalk substituent, if present, is unsubstituted, and more preferably the cycloalk substituent, if present, is either cyclohexyl or adamantyl.

"Longer-length alkyl" refers to an optionally substituted hydrocarbon group joined by single carbon-carbon bonds and which contains 1-20 carbon atoms joined together. Preferably, the longer-length alkyl is 4 to 20 carbon atoms. The longer-length alkyl hydrocarbon group may be straight-chain or contain one or more branches. Longer-length alkyl substituents are each independently selected from the alkenyl substituent list described above. Preferably, the longer-length alkyl is either unsubstituted or has one cycloalk or phenyl substituent. More preferably, the cycloalk substituent, if present, is unsubstituted, and

more preferably the cycloalk substituent, if present, is either cyclohexyl or adamantyl.

"Longer-length alkynyl" refers to an optionally substituted hydrocarbon group containing at least one carbon-carbon triple bond between the carbon atoms, and which contains 2-20 carbon atoms joined together. Preferably, the longer-length alkynyl is 4 to 20 carbon atoms. The longer-length alkynyl hydrocarbon group may be straight-chain or contain one or more branches. Longer-length alkynyl substituents are each independently selected from the alkenyl substituent list described above. Preferably, the longer-length alkynyl is either unsubstituted or has one cycloalk or phenyl substituent. More preferably, the cycloalk substituent, if present, is unsubstituted, and more preferably the cycloalk substituent, if present, is either cyclohexyl or adamantyl.

"Haloalkoxy" refers to oxygen joined to a "haloalk." Preferably, the haloalkoxy is a "lower-haloalkoxy," which is an oxygen joined to a lower-haloalkyl.

20 A. α,α -Disubstituted β -Phenethylamine Derivatives

More preferred calcilytic compounds are Structure I derivatives where R_1 , R_2 , R_3 , R_4 , R_5 , Z , Y_1 and Y_2 are as described above for Structure I α,α -disubstituted arylalkylamine derivatives, including preferred groups (see, 25 Section II, *supra*); and

R_5 is either phenyl substituted with one to four independently selected substituents or an optionally substituted naphthyl having up to four independently selected substituents. R_5 substituents are provided in Section II, 30 *supra.*, including preferred embodiments. More preferably R_5 is either a substituted phenyl comprising a substituent in a

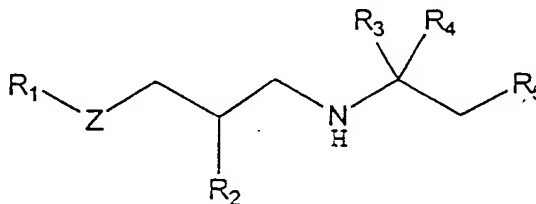
meta or para position, more preferably, the substituent present in a meta or para position is either methyl, ethyl, isopropyl, methoxy, Cl, F, Br, or lower haloalkoxy.

The activity of different calcilytic compounds was measured using the Calcium Receptor Assay described below. Examples of compounds having an $IC_{50} \leq 50 \mu M$ include compounds 1, 9, 17, 25, 29, 42, 56, 79, 90, 101 and 164; examples of preferred compounds having an $IC_{50} < 10 \mu M$ include compounds 2, 3, 7, 8, 26, 27, 32, 33, 35, 37, 39, 41, 45, 48, 49, 59, 61, 66, 68, 71, 75, 93, 98, 103, 104, 110, 111, 114, 123, 124, 125, 128, 132, 144, 147, 152, 155, 158, 161, 162, 169 and 170; and examples of more preferred compounds having an $IC_{50} < 1 \mu M$ include compounds 5, 6, 19, 20, 21, 28, 38, 40, 43, 44, 46, 47, 50, 51, 63, 64, 65, 67, 69, 72, 74, 96, 105, 106, 109, 112, 113, 115, 116, 117, 118, 119, 120, 121, 122, 126, 127, 129, 130, 131, 133, 134, 135, 136, 137, 138, 139, 140, 141, 142, 143, 145, 146, 148, 149, 150, 151, 153, 154, 156, 157, 159, 160, 163, 165, 166, 167, and 168.

B. Structure II Compounds

Structure II compounds have the following structure:

Structure II



In one embodiment R_1 , R_2 , R_3 , and R_4 are as described above for Structure I α, α -disubstituted arylalkylamine

derivatives, including preferred groups (see, Section II, *supra*); and

R_2 is either an optionally substituted naphthyl having one to four substituents independently selected from the group consisting of methyl, ethyl, isopropyl, methoxy, Cl, F, Br, or lower haloalkoxy, preferably the naphthyl is unsubstituted; or a substituted phenyl having one to four substituent with at least one substituent in a *meta* or *para* position selected from the group consisting of: lower alkyl, methoxy, Cl, F, Br, and lower haloalkoxy, more preferably a methoxy is present in the *para* or *meta* position; even more preferably, the remaining R_2 substituents are independently selected from the group consisting of: methoxy, lower-haloalkyl, S-lower alkyl, lower-haloalkoxy, lower alkyl, halogen, SH, CN, OH, Cl, F, and Br.

provided that R_1 is not 6-CN-2-pyridyl; and

further provided that if R_2 is 3,4 dimethoxy-phenyl, then R_1 is not $\text{CH}_3(\text{CH}_2)_5\text{O-phenyl}$; 2-cyclopentyl-phenyl; 2-Cl-phenyl; 2-CN-phenyl; 2-(3-furanyl)phenyl; or 4-(1,2,-benzisothiazol); preferably, R_2 is not 3,4 dimethoxy phenyl;

further provided that if R_2 is 4-methoxy-phenyl, then R_1 is not 2-cyclopentyl-phenyl; 2- CH_3 -phenyl; 2-benzyl-phenyl; 3- CH_3 , 4- CH_3SO_2 -phenyl; or 4-(1,2,-benzisothiazol);

further provided that if R_2 is 4-Cl-phenyl, then R_1 is not 2- CH_3 -phenyl, 5-*iso*-propyl-phenyl; 2- CH_3 -phenyl; 4- CH_3 -phenyl; phenyl; 2-Cl-phenyl; 4-Cl-phenyl; 2-methoxy, 4- CH_3CHCH -phenyl; 3,4 CH_3 -phenyl; 2,4 CH_3 -phenyl; 2,3 CH_3 -phenyl; 2-*iso*-propyl, 5- CH_3 -phenyl; pyridyl; or 1-imidazole; 4-(1,2,-benzisothiazol); preferably, R_1 is either not 4-Cl, or R_1 is 3,4 dichlorophenyl; and

further provided that if R_2 is 3,5, dimethyl, 4-methoxy-phenyl, then R_1 is not 4- CH_3 , 6-CN-2-pyridyl; or

thiophenecarboxamide; preferably, R₅ is not 3,5, dimethyl, 4-methoxy-phenyl.

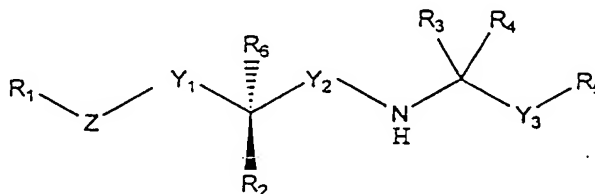
In another embodiment, R₂, R₃, and R₄ are as described above for Structure I α,α -disubstituted arylalkylamine derivatives, including preferred groups (see, Section II, *supra*);

R₅ is either an optionally substituted naphthyl having one to four substituents independently selected from the group consisting of methyl, ethyl, isopropyl, methoxy, Cl, F, Br, and lower haloalkoxy, preferably the naphthyl is unsubstituted; or a substituted phenyl having one to four substituent with at least one substituent in a *meta* or *para* position selected from the group consisting of: methyl, ethyl, isopropyl, methoxy, Cl, F, Br, and lower haloalkoxy, more preferably a methoxy is present in the *para* or *meta* position; even more preferably, the remaining R₅ substituents are independently selected from the group consisting of: methoxy, lower-haloalkyl, S-lower alkyl, lower-haloalkoxy, lower alkyl, halogen, SH, CN, OH, Cl, F, and Br; and

R₁ is either 2-CN-phenyl, 2,3-dichloro phenyl, 2-nitro-phenyl, 2-cyano-3-chloro-phenyl, 2,3-dichloro-4-sulfamoyl-phenyl, an optionally substituted pyridyl, an optionally substituted benzothiopyranyl, or an optionally substituted carbazole, where the optionally present substituents for the pyridyl, benzothiopyranyl, and carbazole as described in Section II *supra*, for aryl R₁ substituents, including preferred substituents, and are even more preferably independently selected from the group consisting of: methoxy, lower-haloalkyl, S-lower alkyl, lower-haloalkoxy, lower alkyl, halogen, SH, CN, OH, Cl, F, Br and sulfamoyl.

C. R₂-group Stereochemistry

The different calcilytic compounds described herein can have different stereochemistry around different groups. In an embodiment of the present invention the Structure I compounds have the following absolute configuration structure with respect to R₂:



III. Pharmaceutical Composition

The calcilytic compounds described herein can be formulated as a pharmaceutical composition to facilitate the administration of the compound to a patient. Preferred formulations contain a pharmaceutically acceptable carrier and a calcilytic compound as described in Section II, *supra.*, including the different embodiments.

Examples of suitable carriers are provided below, in Section V, "Administration," and include calcium carbonate, calcium phosphate, lactose, glucose, sucrose, gelatin, vegetable oils, polyethylene glycols and physiologically compatible solvents.

IV. TREATMENT OF DISEASES OR DISORDERS

Compounds inhibiting calcium receptor activity can be used to confer beneficial effects to patients suffering from a variety of diseases or disorders. Diseases or disorders which can be treated using a calcilytic compound are known in the art and can be identified using the present application as a guide. For example, diseases or disorders can be

identified based on the functional responses of cells regulated by calcium receptor activity.

Diseases and disorders which can be treated using the calcilytic compounds described herein include those due to
5 different cellular defects related to calcium receptor activity in different cells, such as a defective calcium receptor or an abnormal number of calcium receptors, a defective intracellular protein acted on by a calcium receptor, or a defective protein or an abnormal number of
10 proteins acting on a calcium receptor.

Functional responses of cells regulated by the calcium receptor are known in the art, including PTH secretion by parathyroid cells, calcitonin secretion by C-cells, bone resorption by osteoclasts, and Ca^{2+} secretion by kidney cells.
15 Such functional responses are associated with different diseases or disorders.

For example, isolated osteoclasts respond to increases in the concentration of extracellular Ca^{2+} with corresponding increases in $[\text{Ca}^{2+}]_i$ arising partly from the mobilization of
20 intracellular Ca^{2+} . Increases in $[\text{Ca}^{2+}]_i$ in osteoclasts are associated with the inhibition of bone resorption.

Renin secretion from juxtaglomerular cells in the kidney is depressed by increased concentrations of extracellular Ca^{2+} . Extracellular Ca^{2+} causes the mobilization of
25 intracellular Ca^{2+} in these cells. Other kidney cells respond to extracellular Ca^{2+} as follows: elevated Ca^{2+} inhibits formation of $1,25(\text{OH})_2$ -vitamin D by proximal tubule cells, stimulates production of calcium-binding protein in distal tubule cells, and inhibits tubular reabsorption of Ca^{2+} and
30 Mg^{2+} in the thick ascending limb of Henle's loop (MTAL), and reduces vasopressin action in the cortical collecting duct.

Other examples of functional responses affected by extracellular Ca^{2+} include promoting differentiation of

intestinal goblet cells, mammary cells, and skin cells; inhibiting atrial natriuretic peptide secretion from cardiac atria; reducing cAMP accumulation in platelets; altering gastrin and glucagon secretion; acting on perivascular nerves
 5 to modify cell secretion of vasoactive factors; and affecting cells of the central nervous and peripheral nervous systems.

Diseases and disorders which might be treated or prevented, based upon the affected cells, include bone and mineral-related diseases or disorders; hypoparathyroidism;
 10 those of the central nervous system such as seizures, stroke, head trauma, spinal cord injury, hypoxia-induced nerve cell damage, such as occurs in cardiac arrest or neonatal distress, epilepsy, neurodegenerative diseases such as Alzheimer's disease, Huntington's disease and Parkinson's
 15 disease, dementia, muscle tension, depression, anxiety, panic disorder, obsessive-compulsive disorder, post-traumatic stress disorder, schizophrenia, neuroleptic malignant syndrome, and Tourette's syndrome; diseases involving excess water reabsorption by the kidney, such as syndrome of
 20 inappropriate ADH secretion (SIADH), cirrhosis, congestive heart failure, and nephrosis; hypertension; preventing and/or decreasing renal toxicity from cationic antibiotics (e.g., aminoglycoside antibiotics); gut motility disorders such as diarrhea and spastic colon; GI ulcer diseases; GI diseases
 25 with excessive calcium absorption such as sarcoidosis; autoimmune diseases and organ transplant rejection; squamous cell carcinoma; and pancreatitis.

While calcilytic compounds of the present invention will typically be used to treat human patients, they may also be
 30 used to treat similar or identical diseases or disorders in other warm-blooded animal species, such as other primates, farm animals such as swine, cattle, and poultry; and sports animals and pets such as horses, dogs and cats.

Preferably, calcilytic compounds are used in the treatment of bone and mineral-related diseases or disorders. Bone and mineral-related diseases or disorders comprise a diverse class of disorders affecting nearly every major organ system in the body. Examples of bone and mineral-related diseases or disorders include osteosarcoma, periodontal disease, fracture healing, osteoarthritis, rheumatoid arthritis, Paget's disease, humoral hypercalcemia malignancy, and osteoporosis. More preferably, calcilytic compounds are used to treat osteoporosis, a disease characterized by reduced bone density and an increased susceptibility to fractures. Osteoporosis is associated with aging, especially in women.

One way of treating osteoporosis is by altering PTH secretion. PTH can have a catabolic or an anabolic effect on bone. Whether PTH causes a catabolic effect or an anabolic effect seems to depend on how plasma levels of PTH are altered. When plasma levels of PTH are chronically elevated, as in hyperparathyroid states, there is a net loss of bone. In contrast, intermittent increases in plasma PTH levels, as achieved by administration of exogenous hormone, result in new bone formation. Anabolic action of PTH on bone is described, for example, by Dempster et al., *Endocrin. Rev.* 14:690-709, 1993.

As demonstrated by the Examples provided below, calcilytic compounds stimulate secretion of PTH. Such calcilytic compounds can be used to increase bone formation in a patient, for example, by intermittent dosing, thus achieving intermittent increases in the circulating levels of PTH.

V. ADMINISTRATION

The calcilytic compounds described by the present invention can be formulated for a variety of modes of administration, including systemic and topical or localized administration. Techniques and formulations generally may be found in Remington's Pharmaceutical Sciences, 18th ed., Mack Publishing Co., Easton, PA, 1990 (hereby incorporated by reference herein).

Suitable dosage forms, in part, depend upon the use or the route of entry, for example, oral, transdermal, transmucosal, or by injection (parenteral). Such dosage forms should allow the compound to reach a target cell whether the target cell is present in a multicellular host or in culture. For example, pharmacological compounds or compositions injected into the blood stream should be soluble. Other factors are known in the art, and include considerations such as toxicity and dosage forms which retard the compound or composition from exerting its effect.

Compounds can also be formulated as pharmaceutically acceptable salts and complexes thereof. Pharmaceutically acceptable salts are non-toxic salts in the amounts and concentrations at which they are administered. The preparation of such salts can facilitate the pharmacological use by altering the physical characteristics of the compound without preventing it from exerting its physiological effect. Useful alterations in physical properties include lowering the melting point to facilitate transmucosal administration and increasing the solubility to facilitate administering higher concentrations of the drug.

The pharmaceutically acceptable salt of the different compounds may be present as a complex. Examples of complexes include an 8-chlorotheophylline complex (analogous to, e.g.,

dimenhydrinate:diphenhydramine 8-chlorotheophylline (1:1) complex; Dramamine) and various cyclodextrin inclusion complexes.

Pharmaceutically acceptable salts include acid addition salts such as those containing sulfate, hydrochloride, fumarate, maleate, phosphate, sulfamate, acetate, citrate, lactate, tartrate, methanesulfonate, ethanesulfonate, benzenesulfonate, *p*-toluenesulfonate, cyclohexylsulfamate and quinate. Pharmaceutically acceptable salts can be obtained from acids such as hydrochloric acid, maleic acid, sulfuric acid, phosphoric acid, sulfamic acid, acetic acid, citric acid, lactic acid, tartaric acid, malonic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, *p*-toluenesulfonic acid, cyclohexylsulfamic acid, fumaric acid, and quinic acid.

Pharmaceutically acceptable salts also include basic addition salts such as those containing benzathine, chlorprocaine, choline, diethanolamine, ethylenediamine, meglumine, procaine, aluminum, calcium, lithium, magnesium, potassium, sodium, ammonium, alkylamine, and zinc, when acidic functional groups, such as carboxylic acid or phenol are present. For example, see Remington's Pharmaceutical Sciences, 18th ed., Mack Publishing Co., Easton, PA, p. 1445, 1990. Such salts can be prepared using the appropriate corresponding bases.

Pharmaceutically acceptable salts can be prepared by standard techniques. For example, the free-base form of a compound is dissolved in a suitable solvent, such as an aqueous or aqueous-alcohol in solution containing the appropriate acid and then isolated by evaporating the solution. In another example, a salt is prepared by reacting the free base and acid in an organic solvent. (See, e.g., PCT/US92/03736, hereby incorporated by reference herein.)

Carriers or excipients can also be used to facilitate administration of the compound. Examples of carriers include calcium carbonate, calcium phosphate, various sugars such as lactose, glucose, or sucrose, or types of starch, cellulose derivatives, gelatin, vegetable oils, polyethylene glycols and physiologically compatible solvents. Examples of physiologically compatible solvents include sterile solutions of water for injection (WFI), saline solution and dextrose.

The calcilytic compounds can be administered by different routes including intravenous, intraperitoneal, subcutaneous, intramuscular, oral, topical (transdermal), or transmucosal administration. For systemic administration, oral administration is preferred. For oral administration, for example, the compounds can be formulated into conventional oral dosage forms such as capsules, tablets, and liquid preparations such as syrups, elixirs, and concentrated drops.

Alternatively, injection (parenteral administration) may be used, e.g., intramuscular, intravenous, intraperitoneal, and subcutaneous. For injection, the compounds of the invention are formulated in liquid solutions, preferably, in physiologically compatible buffers or solutions, such as saline solution, Hank's solution, or Ringer's solution. In addition, the compounds may be formulated in solid form and redissolved or suspended immediately prior to use. Lyophilized forms can also be produced.

Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, bile salts and fusidic acid derivatives. In addition, detergents may be used to

facilitate permeation. Transmucosal administration, for example, may be through nasal sprays, rectal suppositories, or vaginal suppositories.

For topical administration, the compounds of the invention can be formulated into ointments, salves, gels, or creams, as is generally known in the art.

The amounts of various calcilytic compounds to be administered can be determined by standard procedures taking into account factors such as the compound IC_{50} , EC_{50} , the biological half-life of the compound, the age, size and weight of the patient, and the disease or disorder associated with the patient. The importance of these and other factors to be considered are known to those of ordinary skill in the art. Generally, it is an amount between about 0.1 and 50 mg/kg, preferably 0.01 and 20 mg/kg of the animal to be treated.

VI. Examples

The examples provided below, like the other examples provided herein, are not intended to limit the claimed invention, but rather illustrate different aspects and embodiments of the present invention.

EXAMPLE 1: Calcium Receptor Inhibitor Assay

This example illustrates the use of the Calcium Receptor Inhibitor Assay. Calcilytic activity was measured by determining the IC_{50} of the test compound for blocking increases of intracellular Ca^{2+} elicited by extracellular Ca^{2+} in HEK 293 4.0-7 cells stably expressing the human calcium receptor. HEK 293 4.0-7 cells were constructed as described by Rogers et al., *J. Bone Miner. Res.* 10 Suppl. 1:S483, 1995 (hereby incorporated by reference herein). Intracellular Ca^{2+} increases were elicited by increasing extracellular Ca^{2+} from

1 to 1.75 mM. Intracellular Ca^{2+} was measured using fluo-3, a fluorescent calcium indicator.

The procedure was as follows:

1. Cells were maintained in T-150 flasks in selection
5 media (DMEM supplemented with 10% fetal bovine serum and 200 $\mu\text{g}/\text{mL}$ hygromycin B), under 5% CO_2 :95% air at 37 °C and were grown up to 90% confluency.

2. The medium was decanted and the cell monolayer was washed twice with phosphate-buffered saline (PBS) kept at 37
10 °C. After the second wash, 6 mL of 0.02% EDTA in PBS was added and incubated for 4 minutes at 37 °C. Following the incubation, cells were dispersed by gentle agitation.

3. Cells from 2 or 3 flasks were pooled and pelleted (100 x g). The cellular pellet was resuspended in 10-15 mL
15 of SPF-PCB+ and pelleted again by centrifugation. This washing was done twice.

Sulfate- and phosphate-free parathyroid cell buffer (SPF-PCB) contains 20 mM Na-Hepes, pH 7.4, 126 mM NaCl, 5 mM KCl, and 1 mM MgCl_2 . SPF-PCB was made up and stored at 4 °C.
20 On the day of use, SPF-PCB was supplemented with 1 mg/mL of D-glucose and 1 mM CaCl_2 , and then split into two fractions. To one fraction, bovine serum albumin (BSA; fraction V, ICN) was added at 5 mg/mL (SPF-PCB+). This buffer was used for washing, loading and maintaining the cells. The BSA-free
25 fraction was used for diluting the cells in the cuvette for measurements of fluorescence.

4. The pellet was resuspended in 10 mL of SPF-PCB+ containing 2.2 μM fluo-3 (Molecular Probes) and incubated at room temperature for 35 minutes.

30 5. Following the incubation period, the cells were pelleted by centrifugation. The resulting pellet was washed

with SPF-PCB+. After this washing, cells were resuspended in SPF-PCB+ at a density of $1-2 \times 10^6$ cells/mL.

6. For recording fluorescent signals, 300 μ L of cell suspension were diluted in 1.2 mL of SPF buffer containing 1 mM CaCl_2 and 1 mg/mL of D-glucose. Measurements of fluorescence were performed at 37 °C with constant stirring using a spectrofluorimeter. Excitation and emission wavelengths were measured at 485 and 535 nm, respectively. To calibrate fluorescence signals, digitonin (5 mg/mL in ethanol) was added to obtain F_{max} , and the apparent F_{min} was determined by adding Tris-EGTA (2.5 M Tris-Base, 0.3 M EGTA). The concentration of intracellular calcium was calculated using the following equation:

Intracellular calcium = $(F - F_{\text{min}} / F_{\text{max}}) \times K_d$; where $K_d = 400$ nM.

7. To determine the potential calcilytic activity of test compounds, cells were incubated with test compound (or vehicle as a control) for 90 seconds before increasing the concentration of extracellular Ca^{2+} from 1 to 2 mM. Calcilytic compounds were detected by their ability to block, in a concentration-dependent manner, increases in the concentration of intracellular Ca^{2+} elicited by extracellular Ca^{2+} .

In general, those compounds having lower IC_{50} values in the Calcium Receptor Inhibitor Assay are more preferred compounds. Compounds having an IC_{50} greater than 50 μM were considered to be inactive. Preferred compounds are those having an IC_{50} 10-50 μM , more preferred compounds have an IC_{50} 1-10 μM , and most preferred compounds have an IC_{50} less than 1 μM .

Examples of compounds having an IC_{50} greater than 50 μM include compounds 22, 24, 34, 36, 52, 53, 54, 55, 58, 60, 62, 70, 84, 92, 99, and 102.

EXAMPLE 2: Adrenergic Receptor Activity

Structure I α, α -disubstituted arylalkylamine derivatives include compounds which have both calcilytic activity and β -adrenergic receptor activity. If desired, β -adrenergic activity can be reduced using appropriate functional groups and structural modifications. Modifications which can be carried out to reduce β -adrenergic receptor activity include using alternative R_2 groups and using absolute stereochemistry opposite to that which occurs in active β -adrenergic receptor antagonists, which provides compounds corresponding to the *R* enantiomer when R_2 is OH. β -adrenergic receptor activity and binding to the β -adrenergic receptor can be measured using standard techniques. For example, see Riva et al., *Mol. Pharmacol.* 36:201-210, 1989.

In one embodiment of the present invention the calcilytic compounds have an $IC_{50} \geq 1.0$ nM, at the β -adrenergic receptor as measured using the " β -Adrenergic Receptor Binding Assay" described below. In other embodiments, using the β -Adrenergic Receptor Assay calcilytic compounds have an $IC_{50} \geq 1.0$ μ M, and $IC_{50} \geq 10.0$ μ M.

The " β -Adrenergic Receptor Binding Assay" is carried out as follows: Incubations are performed in polypropylene reaction tubes in a 37 °C water bath. To each tube 50 μ L of test sample is added, followed by 300 μ L of assay buffer (50 mM Tris-HCl, pH 7.5), and 50 μ L of 20 nM [3 H]-dihydroalprenolol. The binding reaction is initiated by the addition of 100 μ L of 3.75 mg/mL well-washed rat cortex membranes in assay buffer, and allowed to incubate at 37 °C for 30 minutes. Non-specific binding is determined in the presence of 10 μ M alprenolol. The final concentration of reactants is: 2 nM [3 H]-dihydroalprenolol, and 75 mg/mL rat cortex membrane in a reaction volume of 0.5 mL.

The binding reaction is terminated by rapid filtration with ice-cold assay buffer onto GF/C filters (Brandel, Gaithersburg, MD) which have been soaked for 15 minutes in assay buffer. The reaction is first diluted with 3 mL of
5 cold assay buffer (4 °C), then aspirated onto the filter followed by 3 x 3 mL washes. Filter disks are placed in 7-mL polypropylene scintillation vials with 5 mL of ScintiSafe 50% (Fisher Scientific, Pittsburgh, PA), and counted overnight.

EXAMPLE 3: Stimulation of PTH Secretion

10 This example illustrates the ability of different calcilytic compounds to exert a biological effect on PTH secretion. PTH secretion was determined using dissociated bovine parathyroid cells as described below for Compounds 32, 33, and 38. Compounds 32, 33, and 38 all stimulated PTH
15 secretion with an EC₅₀ of less than 10 μM.

Stimulation of PTH secretion was assayed as follows:

Preparation of Dissociated Bovine Parathyroid Cells

Parathyroid cell buffer (PCB) contains (mM): NaCl, 126; KCl, 4; MgSO₄, 1; K₂HPO₄/KH₂PO₄, 0.7; Na-Hepes, pH 7.45, and
20 variable amounts of CaCl₂, as specified (reagent grade). PCB was typically supplemented with bovine serum albumin (BSA fraction V; ICN Biomedicals, Inc., Costa Mesa, CA; catalog #81-003) and 1 mg/mL of D-glucose (reagent grade) as indicated. Percoll purification buffer was prepared
25 immediately before use by mixing 8 mL of Percoll (Pharmacia LKB, Alameda, CA; catalog #17-0891-01) and 7 mL of a twice-concentrated PCB solution without phosphate and containing 2 mM CaCl₂.

Parathyroid glands were obtained from calves within
30 minutes of slaughter at an abattoir and shipped via overnight express on ice in PCB containing 1.25 mM CaCl₂. The glands

were trimmed and minced in ice-cold PCB containing 1.25 mM CaCl_2 , 1 mg/mL of D-glucose, and 2% BSA. Dissociated cells were obtained by collagenase digestion by vigorously shaking the minced tissue at 37 °C in PCB containing 0.5 to 1.0%
 5 Collagenase P (Boehringer Mannheim, Indianapolis, IN; catalog #1249 002), 2 to 5 units of deoxyribonuclease (Sigma, St. Louis, MO; catalog #D-0876), 1 mg/mL of D-glucose, and 1.25 mM CaCl_2 (reagent grade). The cell suspension was triturated at 30 minute intervals using 25- and 10-mL pipettes as the
 10 minced tissue was digested and the cells were dispersed.

Cells were pooled at 1-hour intervals by filtering the cell suspension through a 250- μm Nitex screen into 15-mL polystyrene centrifuge tubes and spinning at 100 x g for 5 minutes at 22 °C. The pooled cell pellet was resuspended in
 15 Percoll purification buffer and purified by centrifugation at 14,500 x g for 20 minutes at 4 °C. Dissociated parathyroid cells equilibrated within a buoyant density of 1.048-1.062 above a dense band of red blood cells and below a diffuse band that contains adipocytes, strands of collagen, and
 20 damaged cells.

The dissociated parathyroid cells were removed with a sterile pipette and washed 3 to 4 times under sterile conditions in a 1:1 mixture of Ham's F-12 and Dulbecco's modified Eagle's medium (F-12/DMEM, Sigma, St. Louis, MO; catalog #D 8900) supplemented with 0.5% BSA, 100 U/mL of
 25 penicillin, 100 $\mu\text{g/mL}$ of streptomycin (Gibco BRL, Grand Island, NY; catalog #15140-031), and 20 $\mu\text{g/mL}$ of gentamicin (Sigma, St. Louis, MO; catalog #G 1397).

The cells were finally resuspended in F-12/DMEM
 30 supplemented with lower antibiotic concentrations (10 U/mL of penicillin, 10 $\mu\text{g/mL}$ of streptomycin, and 4 $\mu\text{g/mL}$ of gentamicin). This latter medium lacks serum and contained ITS (insulin, transferrin, selenous acid, BSA, and linoleic

acid; Collaborative Biomedical Products, Bedford, MA; catalog #40352).

Cells were incubated in T-75 flasks at 37 °C in a humid atmosphere of 5% CO₂ in air. Parathyroid cells were collected for use by decanting the flasks after 18 to 24 hours in primary culture. The concentrations of gentamicin and streptomycin used here are considerably below the EC₅₀ for mobilization of intracellular calcium (150 and 600 μM, respectively).

10 Measurement of Parathyroid Hormone (PTH) Secretion

Sulfate, phosphate-free parathyroid cell buffer (SPF-PCB) contains (mM): NaCl, 126; KCl, 5; MgCl₂, 1; Na-Hepes, pH 7.45, and variable amounts of CaCl₂ as specified (reagent grade). SPF-PCB was typically supplemented with bovine serum albumin (BSA fraction V; ICN Biomedicals, Inc., Costa Mesa, CA; catalog #81-003) and 1 mg/mL of D-glucose (reagent grade) as indicated.

Incubations were performed in triplicate in 12 x 75 mm polypropylene or polystyrene tubes to which were added 2.5 μL of test compound. The tubes were kept on ice until the drug additions were completed, then were transferred to a water bath at 37 °C, and the reactions were initiated by the addition of 0.2 mL of a suspension of dissociated cells at a density of 1 to 2 million cells/mL in SPF-PCB containing 0.5 mM Ca²⁺, 1 mg/mL of D-glucose, and 0.1% BSA. Incubation was for 30 minutes and the reaction was terminated by placing the tubes on ice. Cells were pelleted by gentle centrifugation (500 x g for 10 minutes at 4 °C) and 0.1 mL of supernatant was removed and stored at -20 °C.

30 Amino-terminal bovine PTH was measured by radioimmunoassay (RIA) using goat anti-hPTH antiserum H₂, HPLC-purified ¹²⁵I-hPTH (1-34) and bovine PTH (1-84) standards.

Serial dilutions of bPTH standards (1,000 pg/25 μ L to 3.8 pg/25 μ L) were done in 50 mM Tris, pH 7.4, containing 0.5 mM Na azide and 2% bovine serum albumin (diluent). Standards and samples were incubated for 2-3 days at 4 °C in the presence of antiserum after which 1,500-2,000 cpm label/tube was added. After an additional incubation for 1 to 2 days at 4 °C, dextran-coated charcoal was added to separate bound vs. free label. The contents of each tube were mixed and the charcoal was pelleted by centrifugation. The supernatants were decanted into 12 x 75 mm polystyrene tubes and counted in a Packard Cobra gamma counter.

Example 4: General Procedures for the Preparation of Calcilytic Compounds

The calcilytic compounds described by the present invention can be prepared using standard techniques. For example, an overall strategy for preparing preferred compounds described herein can be carried out as described in this section. The examples which follow illustrate the synthesis of specific compounds. Using the protocols described herein as a model, one of ordinary skill in the art can readily produce other Structure I compounds.

All reagents and solvents were obtained from commercial vendors. Starting materials (e.g., amines and epoxides) were synthesized using standard techniques and procedures. GC/EI-MS (Gas Chromatographic/Electron-Impact Mass Spectrometric) analyses were performed on HP-5890 Series II gas chromatographs equipped with HP-Ultra-2 or HP-5MS columns (30 mm x 0.25 mm ID) and HP-5971 or HP-5972 Mass Selective Spectrometric Detectors (MSD's) were used. MPLC (Medium-Pressure Liquid Chromatography) separations were carried out on silica gel (400 mesh) using an FMI pump, ISCO UV-6 detector (254 nm) and FOXY 200 fraction collector. HPLC

(High Performance Liquid Chromatography) was performed using RAININ HP-XL pumps and Dynamix UV-1 detectors (254 nm).

Examples of specific separation conditions and details are given in the individual experimental descriptions

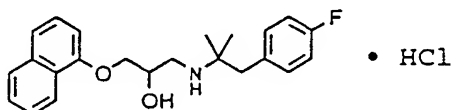
5 provided in the examples below. Chiral HPLC separations were carried out using a Beckman System Gold HPLC and UV detector (254 nm) on Diacel® ChiralCel OD columns (Chiral Technologies, Inc., Exton, PA 19341).

NMR (Nuclear Magnetic Resonance) spectroscopy was
10 performed on a Varian Gemini 300 spectrometer. Proton and carbon spectra were taken at 300 MHz and 75 MHz, respectively. NMR resonances are reported in ppm relative to tetramethylsilane (TMS) with the following descriptors for the observed multiplicities: s (singlet), d (doublet), t
15 (triplet), q (quartet), dd (doublet of doublets) and m (multiplet). J_{AB} coupling constants are reported in Hz. Elemental analyses were performed and FT-IR data were acquired by Oneida Research Services, Inc., Whitesboro, NY 13492.

20 A general procedure used to synthesize many of the compounds was carried out as follows: A solution of glycidyl ether (i.e., 1,2-epoxy-3-phenoxypropane, 1 mmol) and excess amine (typically 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, 1.5 mmol) in absolute ethanol (2 mL) is stirred overnight at
25 50-60 °C. The product is purified by one of three general methods: (1) conversion to the hydrochloride salt, followed by Reversed-Phase High-Performance Liquid Chromatography (RP-HPLC, 0.1% HCl/acetonitrile); (2) conversion to the hydrochloride salt, followed by recrystallization from water-
30 methanol or acetonitrile; and (3) purification by normal-phase chromatography (column chromatography or preparative, thin-layer chromatography (TLC)). Hydrochloride salts were

also prepared by treatment of the corresponding free base in diethyl ether with 1M HCl (in diethyl ether).

EXAMPLE 5: Preparation of N-[2-Hydroxy-3-(1-naphthoxy)propyl]-1,1-dimethyl-2-(4-fluorophenyl)ethylamine Hydrochloride, Compound 2

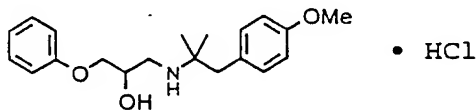


A stirred suspension of sodium hydride (4.0 g of 60% NaH in mineral oil, 100 mmol) in dimethylformamide (DMF, 100 ml) was treated with 1-naphthol (14.42 g, 100 mmol). After stirring for 1 hour at ambient temperature (room temperature), the reaction was treated with epichlorohydrin (10.18 g, 110 mmol) and stirred for 1 hour at 100 °C. The reaction was diluted with water and transferred to a separatory funnel using diethyl ether (500 ml). The organic phase was washed with 10% aqueous NaHCO₃ (3 x 200 ml), dried over anhydrous sodium sulfate, filtered, and concentrated. Kugelrohr distillation (~100 microns) yielded 1-naphthyl glycidyl ether as a clear, colorless oil: GC/EI-MS, *m/z* (rel. int.) 200 (M⁺, 61), 184 (1), 169 (5), 157 (12), 144 (79), 129 (16), 115 (100), 101 (3), 89 (16).

A stirred solution of 1-naphthyl glycidyl ether (400 mg, 2 mmol) and 1,1-dimethyl-2-(4-fluorophenyl)ethylamine (334 mg, 2 mmol) in absolute ethanol (2 mL) was heated at 50-60 °C for 16 hours. Chromatography of the resulting reaction mixture through silica (5 x 30 cm) using a gradient of chloroform to 5% methanol in chloroform afforded the free base of the title compound: GC/EI-MS, *m/z* (rel. int.) 368 (M⁺, 1), 352 (2), 258 (100), 183 (5), 157 (4), 127 (5), 115 (18), 109 (23), 71 (30).

The free base in diethyl ether was treated with excess 1M HCl (diethyl ether). The resulting solid was recrystallized from hot acetonitrile to afford 300 mg of the title compound as a crystalline solid: $^1\text{H-NMR}$ (DMF-D_7) δ 9.9 (1H, br s), 9.5 (1H, br s), 8.33 (1H, d, $J=9$), 7.91 (1H, d, $J=9$), 7.57-7.50 (3H, m), 7.48-7.41 (3H, m), 7.19 (2H, t, $J=10$), 7.03 (1H, d, $J=7$), 6.37 (1H, br d, $J=5$), 4.67 (1H, br s), 4.31 (2H, br t, $J=6$), 3.61 (1H, br t), 3.42 (1H, br t), 3.31 (2H, s), 1.47 (3H, s), 1.46 (3H, s); $^{13}\text{C-NMR}$ (DMF-D_7) δ 161.5, 158.2, 152.2, 132.5, 130.8, 130.7, 129.7, 125.4, 124.4, 124.1, 124.5, 120.0, 118.3, 113.1, 112.8, 103.1, 68.1, 64.2, 57.9, 43.5, 40.3, 20.2.

EXAMPLE 6: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 3



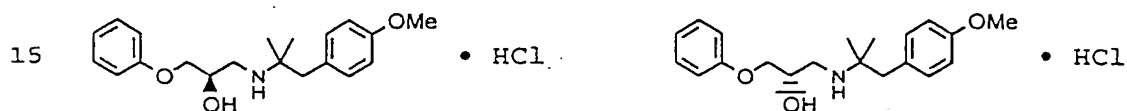
A cooled ($-78\text{ }^{\circ}\text{C}$) solution of diisopropylamine (65 g, 642 mmol) in tetrahydrofuran (THF, 800 mL) was treated with 244 mL of 2.5 M *n*-butyl lithium (610 mmol) in hexane. The reaction was stirred for 30 minutes at room temperature, cooled to $-78\text{ }^{\circ}\text{C}$ and treated dropwise with isobutyric acid (26.8 g, 305 mmol) and hexamethylphosphoramide (HMPA, 54.7 g, 305 mmol). The reaction was stirred for 30 minutes at room temperature and treated with 4-methoxybenzyl chloride (43.4 g, 277 mmol). The reaction was stirred for 48 hours at room temperature and treated with 10% HCl (200 mL). The reaction was concentrated to 300 mL and diluted to 600 mL with water. The resulting solution was extracted with diethyl ether (2 x 300 mL) and the combined ether extracts were washed with 10%

HCl (2 x 200 mL). The ether extract was then extracted with 1N NaOH (3 x 200 mL). The combined 1N NaOH washes were made acidic (pH 1) by the addition of concentrated HCl, and the resulting solution was extracted with diethyl ether (3 x 300 mL). The combined ether extracts were dried over anhydrous magnesium sulfate, filtered, and concentrated to afford 32.6 g of 2,2-dimethyl-3-(4-methoxyphenyl)propionic acid as an oil: GC/EI-MS, m/z (rel. int.) 208 (M^+ , 7), 121 (100), 91 (5), 77 (6).

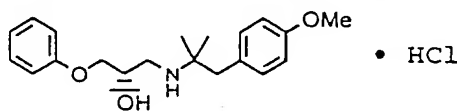
Triethylamine (16.8 g, 166 mmol) and 2,2-dimethyl-3-(4-methoxyphenyl)propionic acid (32.6 g, 157 mmol) were dissolved in 30 mL of water and enough acetone to maintain solubility at 0 °C. A solution of ethyl chloroformate (20.1 g, 185 mmol) in acetone (100 mL) was then added dropwise. An aqueous solution (95 mL) of sodium azide (12.9 g, 198 mmol) was then added dropwise and the resulting reaction mixture stirred 45 minutes at room temperature. The intermediate acyl azide was then extracted into toluene (200 mL). The organic extract was washed with water, dried over anhydrous magnesium sulfate, and heated at 100 °C until the evolution of nitrogen ceased (~45 min). The toluene was removed under vacuum and replaced with benzyl alcohol. The solution was then heated at 100 °C for 16 hours. The excess benzyl alcohol was removed under vacuum. The resulting benzyl carbamate was dissolved in absolute ethanol (200 mL) and reduced in the presence of palladium hydroxide (2 g) under 90 p.s.i. hydrogen for 4 hours at room temperature. The reaction was filtered and concentrated to a yellow oil. Vacuum distillation afforded 13.0 g of 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine as a clear, colorless oil: GC/EI-MS, m/z (rel. int.) 180 ($M+1$, 1), 164 (5), 121 (25), 91 (5), 78 (19), 58 (100).

1,2-Epoxy-3-phenoxypropane (150 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare the free base of the title compound using the method of Example 5, *supra*. The hydrochloride salt
 5 was prepared by dilution of the reaction mixture with HCl (3 mmol) and water. Reversed-phase high-performance liquid chromatography (RP-HPLC, 0.1%/HCl to acetonitrile) of the resulting solution yielded 35 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 314 (M-15, 1), 209
 10 (19), 208 (100), 163 (6), 120 (19), 114 (7), 106 (6), 77 (12), 70 (9), 69 (15), 58 (6).

EXAMPLE 7: Resolution of the Enantiomers (R) or (S)-N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compounds 32 and 33



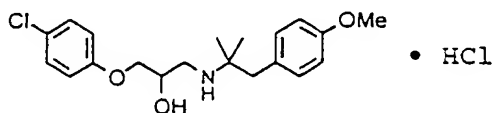
Compound 32



Compound 33

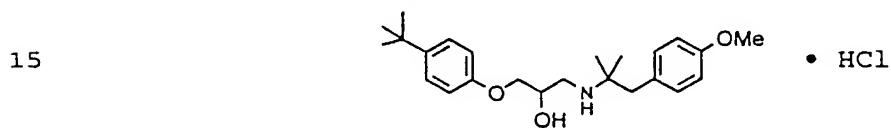
The enantiomers of (R) and (S)-N-(2-hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine hydrochloride (compounds 32 and 33) were obtained by chiral
 20 HPLC of the free base through ChiralCel OD (20 x 2.5 cm) using a combination of hexane-isopropanol containing 0.1% diethylamine (10 mL/min) measuring optical density at 260 nm. GC/EI-MS of each enantiomer gave *m/z* (rel. int.) 330 (M+1, 1), 314 (2), 208 (100), 183 (4), 163 (5), 121 (16), 77 (10),
 25 70 (11). The hydrochloride of each enantiomer was prepared by treatment of the free base in diethyl ether with excess 1M HCl (diethyl ether). Evaporation of the solvent yielded the hydrochloride product as a solid.

EXAMPLE 8: Preparation of N-[2-Hydroxy-3-(4-chlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenylethyl)-amine Hydrochloride, Compound 5



5 Using the method of Example 6, *supra*, 4-chlorophenyl glycidyl ether (185 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare 272 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 348 (M-15, 1), 244 (35), 243 (15), 242
10 (100), 163 (9), 121 (24), 114 (7), 71 (24), 70 (26), 58 (15), 42 (7).

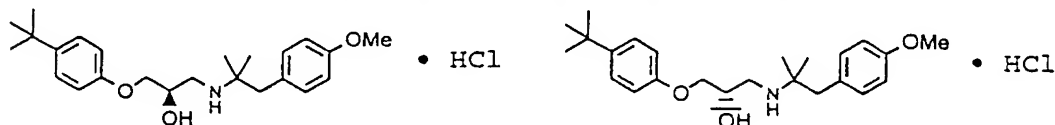
EXAMPLE 9: Preparation of N-[2-Hydroxy-3-(4-*t*-butylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 6



15 Using the method of Example 5, *supra*, 4-*t*-butylphenyl glycidyl ether (206 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare the free base of the title compound. The
20 hydrochloride was prepared by dilution of the reaction mixture with HCl (3 mmol) and water, which caused the product to precipitate. The mixture was heated to effect solution and allowed to cool slowly to crystallize the product. The crystals were collected by filtration, washed with
25 water/MeOH, and dried under vacuum to give 106 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.)

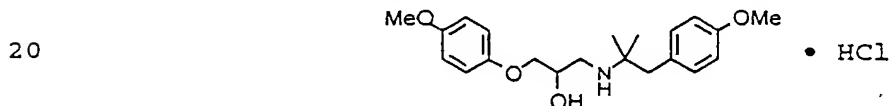
370 (M-15, 0.1), 265 (19), 264 (100), 163 (8), 121 (20), 114 (9), 91 (7), 71 (20), 70 (21), 58 (10), 57 (12).

Example 10: Resolution of the Enantiomers (R) and (S)-N-[2-Hydroxy-3-(4-t-butylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compounds 20 and 21



The enantiomers of (R) and (S)-N-[2-hydroxy-3-(4-t-butylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine hydrochloride were prepared using the method of Example 7, *supra*. GC/EI-MS of each enantiomer gave *m/z* (rel. int.) 386 (M⁺, 1), 370 (2), 264 (100), 163 (10), 135 (4), 121 (36), 91 (8), 70 (11). The hydrochloride salt of each enantiomer was prepared by treatment of the corresponding free base in diethyl ether with excess 1M HCl (diethyl ether). Evaporation of the solvent yielded the corresponding hydrochloride product as a solid.

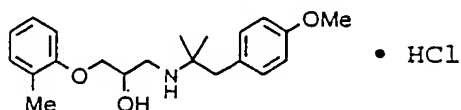
EXAMPLE 11: Preparation of N-[2-Hydroxy-3-(4-methoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 7



Using the method of Example 8, *supra*, 4-methoxyphenyl glycidyl ether (180 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare 231 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 344 (M-15, 0.1), 239 (17), 238 (100), 163

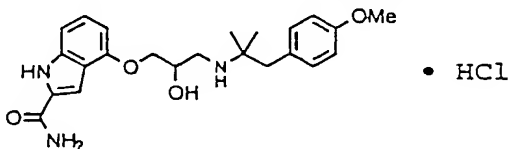
(9), 123 (7), 120 (17), 114 (6), 77 (5), 70 (13), 70 (11), 58 (6).

EXAMPLE 12: Preparation of N-[2-Hydroxy-3-(2-methylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 8



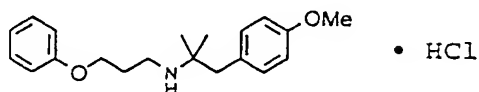
Using the method of Example 9, *supra*, 2-methylphenyl glycidyl ether (164 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare 257 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 328 (M-15, 0.1), 223 (17), 222 (100), 163 (8), 121 (23), 114 (11), 91 (13), 77 (6), 71 (19), 70 (21), 58 (11).

EXAMPLE 13: Preparation of N-[2-Hydroxy-3-(4-(2-carboxamido)indoloxo)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 9



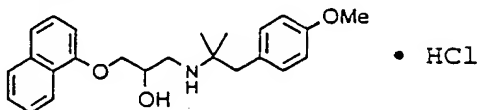
Using the method of Example 6, *supra*, 4-glycidyloxy-2-indolecarboxamide (232 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (269 mg, 1.5 mmol) were used to prepare 222 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 396 (M-15, 0.1), 291 (17), 290 (100), 207 (10), 158 (7), 130 (7), 121 (28), 114 (19), 71 (18), 70 (15).

EXAMPLE 14: Preparation of N-(3-Phenoxypropyl)-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride,
Compound 17



5 To a stirred suspension of 50% KF-Celite (0.35 g, 3
 mmol) in anhydrous acetonitrile (10 mL) was added 1,1-
 dimethyl-2-(4-methoxyphenyl)ethylamine (0.27 g, 1.5 mmol) and
 3-phenoxypropyl bromide (0.484 g, 2.25 mmol). The reaction
 mixture was refluxed under nitrogen for 6 hours, followed by
 10 stirring at room temperature for 62 hours. The mixture was
 filtered and the filtrate, was evaporated. The hydrochloride
 was prepared by dissolving the residue in HCl/methanol. The
 resulting solution was concentrated and dried on a
 lyophilizer. The residue was redissolved in dry methanol and
 15 diluted with diethyl ether, which caused precipitation of 210
 mg of the title compound as a white solid: GC/EI-MS, *m/z*
 (rel. int.) 298 (M-15, 2), 193 (15), 192 (100), 120 (13), 107
 (5), 98 (9), 77 (8), 72 (7), 71 (8), 70 (6), 41 (4).

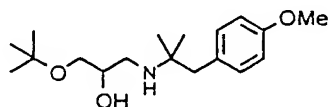
EXAMPLE 15: Preparation of N-[2-Hydroxy-3-(1-
 20 naphthoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
Hydrochloride, Compound 19



Using the method of Example 5, *supra*, 1-naphthyl
 glycidyl ether (1.0 g, 5 mmol) and 1,1-dimethyl-2-(4-
 25 methoxyphenyl)ethylamine (1.0 mg, 5.6 mmol) were used to
 prepare the free base of the title compound. Silica gel
 chromatography of the reaction mixture using 5% methanol in

chloroform afforded 1.66 g (88%) of the purified product:
 GC/EI-MS, m/z (rel. int.) 364 (M-15, 1), 258 (100), 183 (3),
 163 (4), 144 (4), 121 (23), 115 (18), 71 (19): $^1\text{H-NMR}$ (C_6D_6) δ
 8.50 (1H, d, $J=8.0$), 7.65 (1H, d, $J=7.2$), 7.36-7.31 (3H, m),
 5 7.21 (1H, t, $J=7.9$), 6.98 (2H, d, $J=8.6$), 6.71 (2H, d,
 $J=8.6$), 6.62 (1H, d, $J=7.7$), 4.08 (2H, m), 3.93 (1H, m), 3.32
 (3H, s), 2.80 (1H, dd, $J=11.6$ and 3.7), 2.71 (1H, dd, $J=11.4$
 and 6.4), 2.47 (2H, dd, $J=13.3$ and 6.2), 0.94 (3H, s), 0.92
 (3H, s); $^{13}\text{C-NMR}$ (CDCl_3) δ 158.0, 154.3, 134.4, 131.2 (2
 10 carbons), 129.9, 127.4, 126.3, 125.8, 125.2, 121.8, 120.5,
 113.3 (2 carbons), 104.8, 70.4, 68.5, 55.0, 53.2, 46.4, 44.5,
 26.9, 26.8. A portion of the free base in diethyl ether was
 treated with excess 1M HCl (diethyl ether). The resulting
 solid was recrystallized from hot acetonitrile to afford the
 15 title compound as a white solid.

EXAMPLE 16: Preparation of *N*-(2-Hydroxy-3-*t*-
 butoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine,
 Compound 25

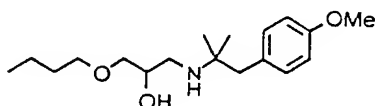


20 Using the method of Example 15, *supra*, *t*-butylglycidyl
 ether (142 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-
 methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to
 prepare 106 mg of the title compound as a clear, colorless
 oil: GC/EI-MS, m/z (rel. int.) 310 (M+1, 0.3), 294 (0.5), 222
 25 (1.8), 188 (67.9), 163 (14.6), 132 (100), 121 (19.1); $^1\text{H-NMR}$
 (CDCl_3) δ 7.09 (2H, d, $J=8.6$), 6.82 (2H, d, $J=8.6$), 3.78 (3H,
 s), 3.68 (1H, m), 3.37 (2H, m), 2.27 (1H, dd, $J=11.5$ and
 4.2), 2.63 (3H, m), 1.19 (9H, s), 1.05 (3H, s), 1.03 (3H, s);

^{13}C -NMR (CDCl_3) δ 156.0, 131.3, 130.3, 113.3, 72.9, 69.7, 64.5, 55.1, 52.9, 46.6, 44.7, 27.4, 26.9, 26.8.

EXAMPLE 17: Preparation of N-(2-Hydroxy-3-butoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

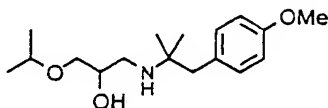
5 Compound 26



Using the method of Example 15, *supra*, *n*-butyl glycidyl ether (143 μL , 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to
 10 prepare 81 mg of the title compound as a clear, colorless oil: GC/EI-MS, m/z (rel. int.) 310 ($M+1$, 0.01), 174 (100), 163 (19), 132 (18), 121 (31), 70 (20); ^1H -NMR (CDCl_3) δ 7.03 (2H, d, $J=8.6$), 6.87 (2H, d, $J=8.6$), 3.74 (1H, m), 3.72 (3H, s), 3.40 (4H, m), 2.73 (3H, m), 2.59 (3H, m), 1.50 (2H, m), 1.30
 15 (2H, m), 1.01 (3H, s), 0.99 (3H, s), 0.87 (3H, t, $J=7.4$); ^{13}C -NMR (CDCl_3) δ 157.9, 131.2, 129.9, 113.2, 73.5, 71.2, 69.0, 54.9, 53.1, 46.2, 44.5, 31.5, 26.6, 26.4, 19.1, 13.8.

EXAMPLE 18: Preparation of N-(2-Hydroxy-3-isopropoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

20 Compound 27

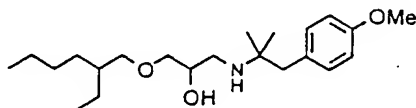


Using the method of Example 15, *supra*, isopropylglycidyl ether (126 μL , 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to
 25 prepare 53 mg of the title compound as a clear, colorless oil; GC/EI-MS, m/z (rel. int.) 296 ($M+1$, 0.2), 280 (1.4),

222 (1.5), 174 (100), 132 (12), 121 (24); $^1\text{H-NMR}$ (CDCl_3) δ 7.03 (2H, d, $J=8.4$), 6.77 (2H, d, $J=8.4$), 3.72 (3H, s), 3.70 (1H, m), 3.53 (1H, m), 3.38 (2H, m), 2.80 (1H, broad s), 2.73 (2H, m), 2.58 (4H, m), 1.09 (6H, m), 1.01 (3H, s), 0.99 (3H, s);
 5 $^{13}\text{C-NMR}$ (CDCl_3) δ 157.9, 131.2, 129.9, 113.2, 73.4, 71.2, 69.0, 54.9, 53.1, 46.2, 44.5, 31.5, 26.6, 26.4, 19.1, 13.8.

EXAMPLE 19: Preparation of *N*-[2-Hydroxy-3-(2-ethyl)hexanoxypopyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine, Compound 28

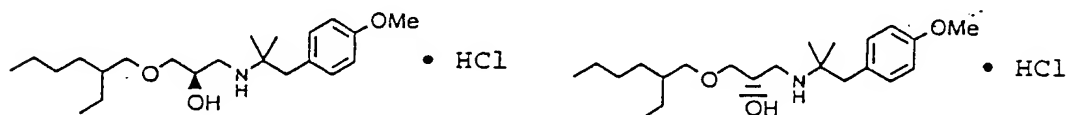
10



Using the method of Example 15, *supra*, 2-ethylhexyl glycidyl ether (209 μL , 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 55 mg of the title compound as a clear, colorless
 15 oil: GC/EI-MS, m/z (rel. int.) 366 ($M+1$, 0.2), 350 (1.1), 244 (100), 222 (2.5), 163 (8.7), 121 (15); $^1\text{H-NMR}$ (CDCl_3) δ 7.06 (2H, d, $J=8.5$), 6.80 (2H, d, $J=8.6$), 3.75 (3H, s), 3.4 (2H, d, $J=5.3$), 3.31 (2H, d, $J=6.0$), 2.88 (1H, broad), 2.78 (1H dd, $J=11.6$ and 4.0), 2.62 (2H, m), 1.46 (1H, q, $J=5.7$), 1.24
 20 (6H, m), 1.03 (4H, m), 0.84 (4H, m); $^{13}\text{C-NMR}$ (CDCl_3) δ 158.0, 131.3, 130.0, 113.3, 74.4, 73.7, 69.0, 55.1, 53.3, 46.3, 44.6, 39.5, 30.5, 29.0, 26.6, 26.4, 23.8, 23.0, 14.0, 11.0; Anal. calculated for $\text{C}_{22}\text{H}_{33}\text{NO}_3$: C, 72.3, H, 10.8, N, 3.8. Found: C, 72.2, H, 9.9, N, 3.6.

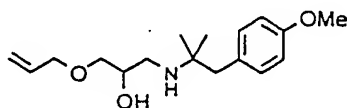
25 EXAMPLE 20: Resolution of the Enantiomers (*R*) and (*S*)-*N*-[2-Hydroxy-3-(2-ethyl)hexanoxypopyl]-1,1-dimethyl-2-

(4-methoxyphenyl)ethylamine Hydrochloride, Compounds 63 and 64



The enantiomers (R) and (S)-N-[2-hydroxy-3-(2-ethyl)hexanoxypentyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine hydrochloride were prepared using the method of Example 7, *supra*. GC/EI-MS of each enantiomer gave *m/z* (rel. int.) 366 (M+1, 1), 350 (2), 244 (100), 222 (3), 163 (12), 133 (9), 121 (21), 115 (11), 100 (4), 71 (21). The hydrochloride salt of each enantiomer was prepared by treatment of the free amine in diethyl ether with excess 1M HCl (diethyl ether). Evaporation of the solvent yielded the hydrochloride product as a solid.

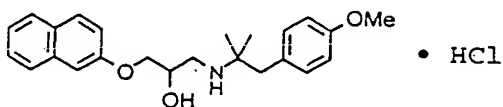
EXAMPLE 21: Preparation of N-(2-Hydroxy-3-allyloxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 29



Using the method of Example 15, *supra*, allyl glycidyl ether (119 μ L, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 79 mg of the title compound as a clear, colorless oil: GC/EI-MS, *m/z* (rel. int.) 294 (M+1, 0.1), 278 (0.9), 222 (1.5), 172 (100), 163 (11), 121 (20); $^1\text{H-NMR}$ (CDCl_3) δ 7.04 (2H, d, $J=8.6$), 6.79 (2H, d, $J=8.6$), 5.85 (1H, ddd, $J=22.2$, 10.5 and 5.7), 5.23 (1H, dd, $J=17.3$ and 1.5), 5.14 (1H, dd, $J=10.3$ and 1.5), 3.96 (2H, d, $J=5.7$), 3.74 (4H, m), 3.43 (2H,

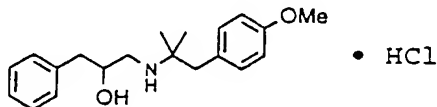
d, $J=5.7$), 2.83 (1H, broad s), 2.77 (1H, dd, $J=11.7$ and 4.1), 2.62 (5H, m), 1.02 (3H, s), 1.01 (3H, s); ^{13}C -NMR (CDCl_3) δ 158.0, 134.5, 131.3, 129.9, 117.0, 113.3, 72.8, 72.2; 69.0, 55.0, 53.3, 46.3, 44.5, 26.6, 26.4.

5 EXAMPLE 22: Preparation of N-[2-Hydroxy-3-(2-naphthoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 35



Using the method of Example 4, *supra*, 2-naphthyl
 10 glycidyl ether (400 mg, 2 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (358 mg, 2 mmol) were used to prepare the free base of the title compound: GC/EI-MS, m/z (rel. int.) 364 (M-15, 1), 258 (100), 183 (2), 163 (3), 144 (4), 127 (10), 121 (22), 115 (20), 71 (11). The free base
 15 in diethyl ether was treated with excess 1M HCl (diethyl ether). The resulting solid was recrystallized from hot acetonitrile to afford 496 mg of the hydrochloride product as a white solid.

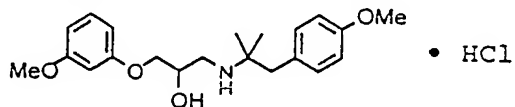
20 EXAMPLE 23: Preparation of N-(2-Hydroxy-3-phenylpropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 37



Using the method of Example 6, *supra*, 2,3-epoxypropylbenzene (1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.25 mmol) yielded 179 mg of the
 25

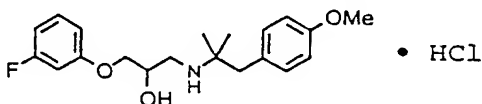
title compound as a white solid: GC/EI-MS, m/z (rel. int.) 298 (M-15, 1), 193 (16), 192 (100), 163 (7), 121 (18), 117 (12), 91 (32), 77 (5), 76 (5), 70 (16), 58 (9).

Example 24: Preparation of *N*-[2-Hydroxy-3-(3-methoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 38



Using the method of Example 6, *supra*, 3-methoxyphenyl glycidyl ether (1.5 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.9 mmol) yielded 403 mg of the title compound as a white solid: GC/EI-MS, m/z (rel. int.) 344 (M-15, 1), 239 (21), 238 (100), 163 (10), 121 (16), 114 (9), 106 (3), 77 (5), 71 (7), 70 (10), 58 (4).

EXAMPLE 25: Preparation of *N*-[2-Hydroxy-3-(3-fluorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 39

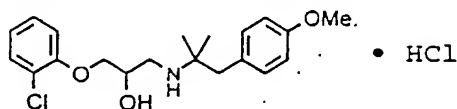


A solution of 3-fluorophenol (1.8 g, 16.1 mmol) in acetone (100 mL) was treated with potassium carbonate (6.65 g, 48.2 mmol) and refluxed under nitrogen for 15 minutes. Epibromohydrin (4.4 g, 32.1 mmol) was then added by syringe, and the mixture was refluxed 3 hours. The mixture was cooled and filtered, and the filtrate evaporated to dryness. The residue was partitioned between ether/water, and the layers separated. The ether layer was washed with saturated NaCl, dried over sodium sulfate and evaporated. The resulting oil

was distilled under vacuum to give 1.2 g of 3-fluorophenyl glycidyl ether.

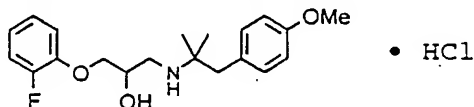
Using the method of Example 6, *supra*, 3-fluorophenyl glycidyl ether (1.5 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.9 mmol) yielded 398 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 332 (M-15, 1), 227 (22), 226 (100), 163 (7), 151 (6), 120 (22), 114 (6), 94 (7), 71 (11), 70 (16), 57 (8).

EXAMPLE 26: Preparation of N-[2-Hydroxy-3-(2-chlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 40



Using the method of Example 25, *supra*, 2-chlorophenyl glycidyl ether (1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.25 mmol) yielded 279 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 348 (M-15, 5), 245 (21), 244 (100), 242 (100), 163 (29), 121 (82), 114 (24), 77 (21), 71 (44), 70 (56), 58 (24).

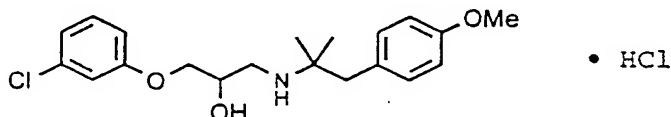
EXAMPLE 27: Preparation of N-[2-Hydroxy-3-(2-fluorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 41



Using the method of Example 25, *supra*, 2-fluorophenyl glycidyl ether (1.5 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.9 mmol) yielded 385 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.)

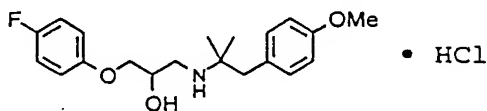
332 (M-15, 2); 227 (20), 226 (100), 163 (4), 125 (3), 121 (15), 78 (4), 77 (4), 71 (7), 70 (9), 58 (3).

EXAMPLE 28: Preparation of N-[2-Hydroxy-3-(3-chlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 43



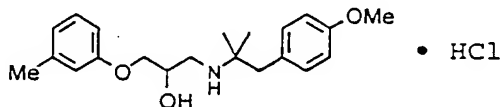
Using the method of Example 25, *supra*, 3-chlorophenyl glycidyl ether (1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.25 mmol) yielded 168 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 348 (M-15, 0.9), 245 (7), 244 (35), 243 (25), 242 (100), 163 (7), 121 (22), 71 (11), 70 (16).

EXAMPLE 29: Preparation of N-[2-Hydroxy-3-(4-fluorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 44



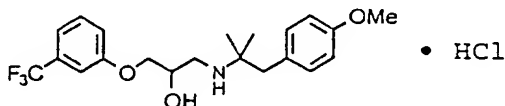
Using the method of Example 25, *supra*, 4-fluorophenyl glycidyl ether (1.5 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.9 mmol) yielded 398 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 332 (M-15, 1), 227 (20), 226 (100), 163 (5), 125 (4), 121 (15), 114 (3), 95 (4), 71 (8), 70 (10), 58 (5).

EXAMPLE 30: Preparation of N-[2-Hydroxy-3-(3-methylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 45



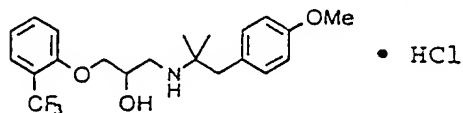
5 Using the method of Example 25, *supra*, 3-methylphenyl glycidyl ether (2.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (2.5 mmol) yielded 400 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 328 (M-15,1), 223 (16), 222 (100), 163 (5), 147 (5), 121
10 (18), 114 (6), 91 (8), 76 (4), 71 (6), 70 (11).

EXAMPLE 31: Preparation of N-[2-Hydroxy-3-(3-trifluoromethylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 46



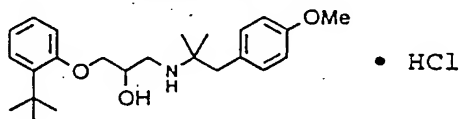
15 Using the method of Example 25, *supra*, 3-trifluoromethylphenyl glycidyl ether (2.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (2.5 mmol) yielded 600 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 382 (M-15, 1), 277 (16), 276 (100), 163 (7), 126
20 (4), 121 (18), 114 (5), 96 (6), 71 (8), 70 (15), 57 (4).

EXAMPLE 32: Preparation of N-[2-Hydroxy-3-(2-trifluoromethylphenoxy)propyl]-1,1-dimethyl-2-(4-

methoxyphenyl)ethylamine Hydrochloride, Compound 47

Using the method of Example 25, *supra*, 2-tri-
fluoromethylphenyl glycidyl ether (2.0 mmol) and 1,1-
5 dimethyl-2-(4-methoxyphenyl)ethylamine (2.5 mmol) yielded 690
mg of the title compound as a white solid: GC/EI-MS, *m/z*
(rel. int.) 382 (M-15, 1), 277 (16), 276 (100), 163 (10), 121
(22), 114 (8), 96 (11), 71 (17), 70 (33), 58 (9), 42 (6).

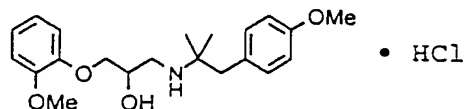
10 EXAMPLE 33: Preparation of N-[2-Hydroxy-3-
(2-*t*-butylphenoxy)propyl]-1,1-dimethyl-2-(4-
methoxyphenyl)ethylamine Hydrochloride, Compound 48



Using the method of Example 25, *supra*, 2-*t*-butylphenyl
glycidyl ether (2.0 mmol) and 1,1-dimethyl-2-(4-
15 methoxyphenyl)ethylamine (2.5 mmol) yielded 540 mg of the
title compound as a white solid: GC/EI-MS, *m/z* (rel. int.)
370 (M-15, 1), 265 (19), 264 (100), 163 (5), 121 (17), 114
(6), 91 (8), 77 (3), 71 (9), 70 (8), 58 (3).

20 EXAMPLE 34: Preparation N-[2-Hydroxy-3-
(2-methoxyphenoxy)propyl]-1,1-dimethyl-2-(4-

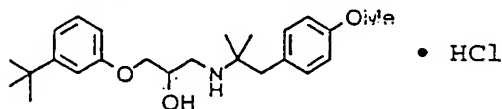
methoxyphenyl)ethylamine Hydrochloride, Compound 49



Using the method of Example 25, *supra*, 2-methoxyphenyl glycidyl ether (2.0 mmol) and 1,1-dimethyl-2-(4-

5 methoxyphenyl)ethylamine (2.5 mmol) yielded 60 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 344 (M-15, 0.1), 239 (15), 238 (100), 163 (9), 122 (7), 121 (20), 114 (13), 77 (10), 71 (19), 70 (21), 58 (7).

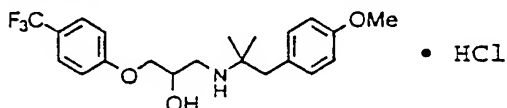
EXAMPLE 35: Preparation of N-[2-Hydroxy-3-(3-
10 *t*-butylphenoxy)propyl]-1,1-dimethyl-2-(4-
methoxyphenyl)ethylamine Hydrochloride, Compound 50



Using the method of Example 25, *supra*, 3-*t*-butylphenyl glycidyl ether (2.0 mmol) and 1,1-dimethyl-2-(4-

15 methoxyphenyl)ethylamine (2.5 mmol) yielded 400 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 370 (M-15, 1), 265 (19), 264 (100), 163 (5), 121 (15), 114 (5), 110 (3), 91 (4), 71 (6), 70 (9), 57 (3).

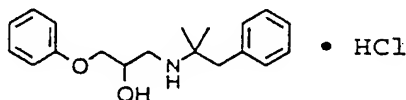
EXAMPLE 36: Preparation of N-[2-Hydroxy-3-(4-
20 trifluoromethylphenoxy)propyl]-1,1-dimethyl-2-(4-
methoxyphenyl)ethylamine Hydrochloride, Compound 51



Using the method of Example 25, *supra*, 4-tri-fluoromethylphenyl glycidyl ether (1.43 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.8 mmol) yielded 270 mg of the title compound as a white solid: GC/EI-MS, m/z (rel. int.) 382 (M-15, 3), 277 (35), 276 (100), 175 (8), 163 (8), 145 (16), 121 (34), 78 (9), 71 (15), 70 (19), 58 (8).

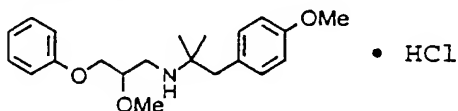
EXAMPLE 37: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-phenylethylamine Hydrochloride, Compound 56

10



Using the method of Example 5, *supra*, 1,2-epoxy-3-phenoxypropane (600 mg, 4 mmol) and 1,1-dimethyl-2-phenylethylamine (596 mg, 4 mmol) yielded the title compound: GC/EI-MS, m/z (rel. int.) 284 (M+1, 1), 208 (100), 162 (1), 133 (7), 91 (27), 77 (15), 70 (22). The free base in diethyl ether was treated with excess 1M HCl (diethyl ether). The resulting solid was recrystallized from hot acetonitrile to afford 596 mg of the hydrochloride product as a white solid.

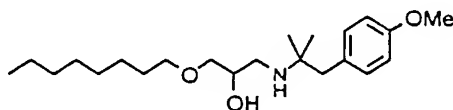
20 EXAMPLE 38: Preparation of N-(2-Methoxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 59



25 Allyl phenyl ether (1.34 g, 10 mmol) and N-bromosuccinimide (1.78 g, 10 mmol) were dissolved in 50 mL of methanol and stirred at room temperature for two days. The

product, a 1:1 mixture of 2-bromo-1-methoxy-3-phenoxypropane and 1-bromo-2-methoxy-3-phenoxypropane, was isolated by evaporating the methanol and dissolving the residue in heptane/ether/water. The organic layer was washed first with water, then brine, dried over sodium sulfate, and evaporated to dryness. The crude mixture (1.47 g, 6 mmol) was dissolved in 6 mL of acetonitrile, to which was added 50% KF-Celite (0.7 g, 12 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine (0.54 g, 3 mmole). The mixture was refluxed under nitrogen for 48 hours, and then cooled and filtered. The filtrate was evaporated to dryness, and the residue was taken up in water and ether. The ether layer was dried over sodium sulfate and concentrated to dryness. The residue was dissolved in 10 mL of diethyl ether and precipitated as the HCl salt by the addition of 10 mL of 1M HCl (diethyl ether). The collected solid was purified by RP-HPLC to yield 330 mg of the title compound as a white solid: GC/EI-MS, m/z (rel. int.) 328 (M-15, 5), 223 (43), 222 (100), 163 (9), 133 (13), 121 (42), 107 (13), 78 (11), 77 (23), 71 (12), 70 (32).

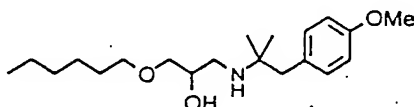
EXAMPLE 39: Preparation of N-(2-Hydroxy-3-octanoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.
Compound 65



Using the method of Example 5, *supra*, 1-octyl glycidyl ether (187 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 105 mg of the title compound as a clear, colorless oil: GC/EI-MS, m/z (rel. int.) 366 (M+1, 0.08), 350 (0.08), 244 (100), 222 (5.8), 163 (12), 121 (18): $^1\text{H-NMR}$ (CDCl_3) δ 7.08 (2H, d,

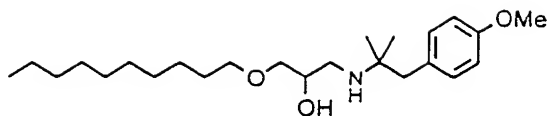
$J=8.6$), 6.82 (2H, d, $J=8.6$), 3.79 (1H, m), 3.77 (3H, s), 3.45 (4H, m), 2.92 (1H, broad s), 2.79 (1H, dd, $J=11.6$ and 4.1), 2.65 (2H, m), 2.55 (2H, m), 1.27 (8H, m), 1.06 (3H, s), 1.04 (3H, s), 0.88 (3H, t, $J=6.7$); ^{13}C -NMR (CDCl_3) δ 158.0, 131.2, 129.9, 113.2, 73.4, 71.6, 68.9, 55.0, 53.3, 46.2, 44.6, 31.7, 29.5, 29.3, 29.1, 26.5, 26.4, 26.0, 22.5, 14.0; FT-IR (film) cm^{-1} 3409 (broad), 1611, 1512, 1245, 1120.

EXAMPLE 40: Preparation of *N*-(2-Hydroxy-3-hexanoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.
 10 Compound 66



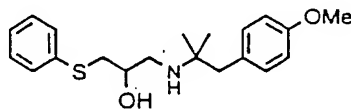
Using the method of Example 5, *supra*, 1-hexyl glycidyl ether (175 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to
 15 prepare 95 mg of the title compound as a clear, colorless oil: GC/EI-MS, m/z (rel. int.) 216 (M-121), 163 (11), 121 (22), 114 (14); ^1H -NMR (CDCl_3) δ 7.05 (2H, d, $J=8.7$), 6.79 (2H, d, $J=8.7$), 3.77 (1H, m), 3.75 (3H, s), 3.41 (4H, m), 2.97 (2H, broad), 2.79 (1H, dd, $J=11.7$ and 4.1), 2.64 (3H, m), 1.52 (2H, m), 1.26 (6H, m), 1.04 (3H, s), 1.03 (3H, s), 0.85 (3H, t, $J=6.7$); ^{13}C -NMR (CDCl_3) δ 158.1, 131.3, 129.9, 113.4, 73.4, 71.7, 68.9, 55.1, 53.6, 46.3, 44.6, 31.6, 29.5, 26.5, 26.4, 25.7, 22.6, 14.0; FT-IR, cm^{-1} 3405 (broad), 1611, 1512, 1245, 1116, 824.

25 EXAMPLE 41: Preparation of *N*-(2-Hydroxy-3-decanoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.
Compound 67



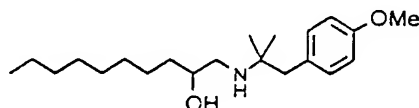
Using the method of Example 5, *supra*, 1-decyl glycidyl ether (235 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 175 mg of the title compound as a clear, colorless oil: GC/EI-MS, m/z (rel. int.) 394 ($M+1$, 1), 378 (6), 273 (97), 272 (100), 222 (9), 163 (37), 121 (61); $^1\text{H-NMR}$ (CDCl_3) δ 7.49 (2H, d, $J=8.5$), 6.82 (2H, d, $J=8.5$ Hz), 3.78 (3H, s), 3.75 (1H, m), 3.45 (4H, m), 2.80 (1H, dd, $J=11.7$ and 4.0), 2.77 (1H, broad s), 2.64 (4H, m), 1.56 (2H, m), 1.26 (16 H, m), 1.06 (3H, s), 1.05 (3H, s), 0.88 (3H, t, $J=6.1$); $^{13}\text{C-NMR}$ (CDCl_3) δ 158.1, 131.3, 130.0, 113.4, 73.5, 71.7, 69.1, 55.1, 53.3, 46.4, 44.6, 31.8, 29.6, 29.5, 29.4, 29.3, 26.7, 26.6, 26.1, 22.6, 14.1; FT-IR (film) cm^{-1} 3115 (broad s), 1612, 1512, 1245, 1121, 824.

EXAMPLE 42: Preparation of N-(2-Hydroxy-3-thiophenylpropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 68



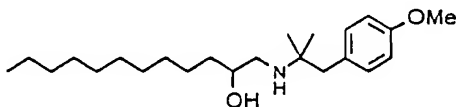
Using the method of Example 25, *supra*, phenyl glycidyl sulfide (3.9 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (4.9 mmol) yielded 194 mg of the title compound as a white solid: GC/EI-MS, m/z (rel. int.) 330 ($M-15$, 4), 226 (21), 225 (58), 224 (100), 163 (25), 149 (25), 123 (100), 121 (75), 77 (18), 71 (22), 70 (26).

EXAMPLE 43: Preparation of N-(2-Hydroxydecanyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 69



Using the method of Example 5, *supra*, 1,2-epoxydecane
 5 (204 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)-
 ethylamine (197 mg, 1.1 mmol) were used to prepare 73 mg of
 the title compound as a clear, colorless oil: GC/EI-MS, *m/z*
 (rel. int.) 214 (M-121, 100), 196 (27.8), 163 (10), 121
 (2.2); ¹H-NMR (CDCl₃) δ 7.08 (2H, d, *J*=8.6), 6.83 (2H, d,
 10 *J*=8.6), 3.79 (3H, s), 3.53 (1H, m), 2.79 (1H, dd, *J*=11.7 and
 2.9), 2.67 (2H, s), 2.42 (1H, dd, *J*=12.6 and 9.6), 1.26 (18
 H, m), 1.08 (3H, s), 0.88 (3H, t, *J*=6.6); ¹³C-NMR (CDCl₃) δ
 158.2, 131.4, 129.8, 113.5, 77.2, 69.8, 55.2, 53.8, 47.8,
 46.5, 35.1, 31.9, 29.8, 29.7, 29.6, 29.3, 26.6, 26.4, 25.7,
 15 22.7, 14.1; FT-IR (film) cm⁻¹ 3399 (broad s), 1612, 1512,
 1246, 1039, 823.

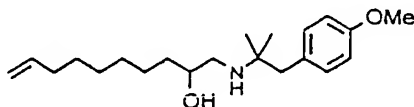
EXAMPLE 44: Preparation of N-(2-Hydroxydodecanyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 70



20 Using the method of Example 5, *supra*, 1,2-epoxydodecane
 (240 μL, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)-
 ethylamine (197 mg, 1.1 mmol) were used to prepare 68 mg of
 the title compound as a clear, colorless oil: GC/EI-MS, *m/z*
 (rel. int.) 363 (M+1, 0.2), 348 (2), 242 (100), 224 (8), 163
 25 (7), 121 (20); ¹H-NMR (CDCl₃) δ 7.08 (2H, d, *J*=8.4), 6.83 (2H,
 d, *J*=8.4), 3.79 (3H, s), 3.50 (1H, m), 2.78 (1H, dd, *J*=11.8
 and 3.1), 2.66 (2H, s), 2.41 (1H, dd, *J*=11.5 and 9.3), 1.26

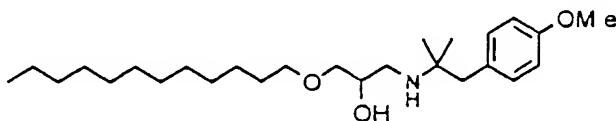
(18 H, m), 1.07 (6H, s), 0.88 (3H, t, $J=6.5$); ^{13}C -NMR (CDCl_3) δ 158.2, 131.4, 129.9, 113.4, 76.9, 70.0, 55.2, 53.6, 47.8, 46.6, 35.1, 31.9, 29.7, 29.6, 29.3, 26.7, 26.5, 25.7, 22.7, 14.1; FT-IR (film) cm^{-1} 3386, 1612, 1512, 1246, 1039, 824.

5 EXAMPLE 45: Preparation of N-(2-Hydroxydec-9-enyl)-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 71



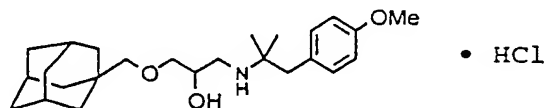
Using the method of Example 5, *supra*, 1,2-epoxy-9-decene
 (202 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)-
 10 ethylamine (197 mg, 1.1 mmol) were used to prepare 80 mg of
 the title compound as a clear, colorless oil: GC/EI-MS, m/z
 (rel. int.) 334 ($M+1$, 0.02), 318 (0.7), 212 (100), 194 (13),
 163 (11), 121 (23); ^1H -NMR (CDCl_3) δ 7.06 (2H, d, $J=8.6$), 6.80
 (2H, d, $J=8.6$), 5.79 (1H, dddd, $J=23.1$, 10.2, 6.6 and 6.6),
 15 4.95 (2H, m), 3.77 (3H, s), 3.52 (1H, m), 2.76 (1H, dd,
 $J=11.7$ and 3.0), 2.64 (1H, s), 2.39 (1H, dd, $J=11.6$ and 9.4),
 2.03 (2H, m), 1.33 (10H, m), 1.05 (6H, s); ^{13}C -NMR (CDCl_3) δ
 158.1, 139.1, 131.4, 129.8, 114.1, 113.4, 69.8, 55.2, 53.7,
 47.8, 46.5, 35.1, 33.8, 29.6, 29.0, 28.8, 26.6, 26.4, 25.7;
 20 FT-IR (film) cm^{-1} 3387 (broad, s), 1612, 1512, 1246, 1038,
 910, 760.

25 EXAMPLE 46: Preparation of N-(3-Dodecanoxy-2-
hydroxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine,
Compound 72



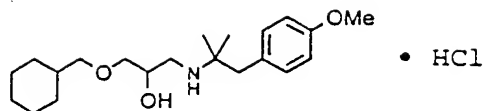
Using the method of Example 5, *supra*, dodecyl glycidyl ether (242 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 121 mg of the title compound as a clear, colorless oil: GC/EI-MS, *m/z* (rel. int.) 422 (*M*+1, 1), 406 (4), 300 (100), 222 (11), 163 (23), 121 (34); ¹H-NMR (CDCl₃) δ 7.09 (2H, d, *J*=8.6), 6.83 (2H, d, *J*=8.6), 3.79 (3H, s), 3.76 (1H, m), 3.45 (4H, m), 2.81 (1H, dd, *J*=7.6 and 4.0), 2.65 (4H, m), 1.53 (2H, m), 1.26 (20H, m), 1.06 (3H, s), 1.05 (3H, s), 0.88 (3H, t, *J*=6.4); ¹³C-NMR (CDCl₃) δ 158.1, 131.3, 130.0, 113.4, 73.5, 71.7, 69.0, 55.1, 53.4, 46.4, 44.6, 31.9, 29.6, 29.4, 29.3, 26.7, 26.6, 26.1, 22.7, 14.1; FT-IR (film) cm⁻¹ 3415 (broad, s), 1612, 1512, 1246, 1121, 825.

EXAMPLE 47: Preparation of N-[2-Hydroxy-3-(1-adamantylmethoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 74



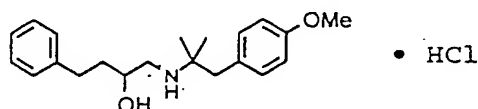
Using the method of Example 9, *supra*, 1,2-epoxy-3-(1-adamantylmethoxy)propane (410 mg, 1.8 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (403 mg, 2.25 mmol) were used to prepare 625 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 386 (*M*-15, 5), 281 (97), 280 (100), 163 (26), 149 (77), 135 (23), 121 (63), 107 (18), 93 (29), 79 (20), 71 (26).

EXAMPLE 48: Preparation of N-(2-Hydroxy-3-cyclohexylmethoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 75



Using the method of Example 6, *supra*, 1,2-epoxy-3-cyclohexylmethoxypropane (212 mg, 1.2 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (279 mg, 1.6 mmol) were used to
 5 prepare 200 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 334 (M-15, .1), 229 (15), 228 (100), 163 (9), 132 (5), 121 (16), 114 (9), 97 (7), 71 (8), 70 (9), 55 (16).

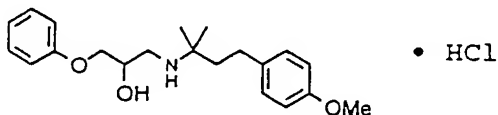
EXAMPLE 49: Preparation of N-(2-Hydroxy-4-phenylbutyl)-
 10 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride.
Compound 79



A solution of *m*-chloroperbenzoic acid (43.5 g, 151 mmol) in chloroform (250 ml) was treated with 4-phenyl-1-butene (20
 15 g, 151 mmol). The reaction was stirred for 1 hour at room temperature and washed with sodium bicarbonate, sodium sulfite, and saturated sodium chloride. The solution was dried over sodium sulfate and evaporated to dryness to afford 3,4-epoxybutylbenzene (100%).

20 Using the method of Example 6, *supra*, 3,4-epoxybutylbenzene (1.4 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.7 mmol) were used to prepare 235 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 312 (M-15, 0.1), 207 (16), 206 (100), 163 (7),
 25 131 (28), 121 (19), 91 (28), 77 (7), 71 (7), 70 (10), 58 (10).

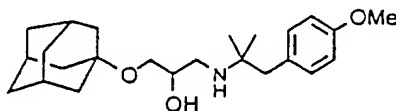
EXAMPLE 50: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-3-(4-methoxyphenyl)propylamine Hydrochloride, Compound 90



5 4-Methoxycinnamonitrile was hydrogenated in ethanol with palladium hydroxide on carbon to give 3-(4-methoxyphenyl)-propionitrile. A mixture of anhydrous cerium (III) chloride (1.99 g, 8.1 mmol) in dry THF (12 mL) was stirred for 3 hours at room temperature, cooled to -78 °C, and treated with MeLi
10 (5.8 mL, 8.1 mmol). After stirring for 1 hour at -78 °C the reaction mixture was treated with 3-(4-methoxy-phenyl)propionitrile (0.45 g, 2.8 mmol). The reaction mixture was stirred for 5 hours at -78 °C and then quenched with ammonium hydroxide. After warming to room temperature,
15 the mixture was filtered, and the filtrate diluted with water and extracted with diethyl ether. The diethyl ether layer was dried over sodium sulfate and evaporated. The crude oil was purified by normal-phase chromatography to give 150 mg of 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine as a light yellow
20 oil.

 Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.62 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.78 mmol) were used to prepare 105 mg of the title compound as a white solid: GC/EI-MS, *m/z*
25 (rel. int.) 343 (M⁺, 4), 209 (14), 208 (99), 161 (13), 122 (9), 121 (100), 77 (17), 72 (25), 71 (12), 70 (18), 58 (13).

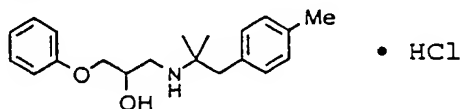
EXAMPLE 51: Preparation of N-[2-Hydroxy-3-(1-adamantanoxyl)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, Compound 96



5 Using the method of Example 16, *supra*, 1-adamantyl glycidyl ether (350 mg, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (197 mg, 1.1 mmol) were used to prepare 207 mg of the title compound as a clear, colorless oil: GC/EI-MS, *m/z* (rel. int.) 338 (M+1, 0.1), 372 (0.3), 266
 10 (65), 163 (1), 135 (100), 121 (16); ¹H-NMR (CDCl₃) δ 7.02 (2H, d, *J*=8.3), 6.74 (2H, d, *J*=8.6), 3.70 (3H, s), 3.64 (1H, m), 3.38 (4H, m), 2.71 (1H, dd, *J*=11.5 and 4.0), 2.57 (3H, m), 2.06 (3H, broad s), 1.64 (6H, broad s), 1.53 (6H, m), 1.13 (4H, apparent t, *J*=6.9), 0.98 (3H, s), 0.97 (3H, s); ¹³C-NMR
 15 (CDCl₃) δ 157.8, 131.2, 130.0, 113.1, 77.8, 70.5, 69.4, 54.9, 52.8, 46.3, 44.6, 32.0, 26.7, 26.6, 25.6, 23.9.

EXAMPLE 52: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methylphenyl)ethylamine Hydrochloride, Compound 98

20



A solution of 2,4,6-triphenylpyrylium tetrafluoroborate (2.97 g, 7.5 mmol) in ethanol (15 mL) was treated with 4-methylbenzylamine (1 g, 8.25 mmol). The reaction was stirred overnight at room temperature and diluted with diethyl ether
 25 to precipitate the product. The product was recrystallized from ethanol/diethyl ether to give 3.15 g of N-(4-

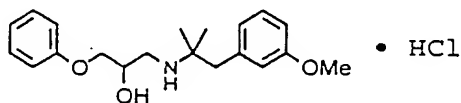
methylbenzyl)-2,4,6-triphenylpyridinium tetrafluoroborate as a tan solid.

Sodium hydride (0.92 g, 60% oil dispersion, 22.9 mmol) was added to methanol (10 mL) at 0 °C, followed by the addition of 2-nitropropane (2.04 g, 22.9 mmol). The reaction mixture was stirred for 30 minutes at room temperature and the methanol was evaporated at reduced pressure. A solution of the *N*-(4-methylbenzyl)-2,4,6-triphenylpyridinium tetrafluoroborate (3.15 g, 7.6 mmol) in DMSO (25 mL) was then added to the dry sodium salt of 2-nitropropane. The mixture was stirred at 60 °C overnight under nitrogen. The reaction was diluted with water, and the product extracted into diethyl ether. The ether layer was washed with saturated NaCl and dried over sodium sulfate. The ether solution was treated with Amberlyst 15 ion-exchange resin to absorb the 2,4,6-triphenylpyridine. The resin was filtered and the filtrate evaporated to yield 1.3 g of pure 1-(4-methylphenyl)-2-methyl-2-nitropropane.

The 1-(4-methylphenyl)-2-methyl-2-nitropropane (1.3 g) was hydrogenated for 5 hours at 65 p.s.i. hydrogen in ethanol (30 mL) using 1.4 g of Raney nickel as catalyst. Removal of the catalyst by filtration and evaporation of the solvent yielded 1.15 g of 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine as a clear oil.

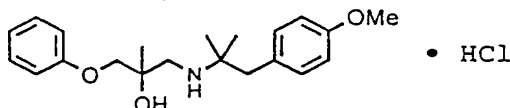
Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (1.3 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.6 mmol) were used to prepare 85 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 298 (M-15, 7), 209 (47), 208 (100), 114 (14), 107 (13), 105 (46), 79 (12), 77 (28), 71 (18), 70 (31), 58 (13).

EXAMPLE 53: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(3-methoxyphenyl)ethylamine Hydrochloride, Compound 99



5 Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (200 mg, 1.3 mmol) and 1,1-dimethyl-2-(3-methoxyphenyl)ethylamine (263 mg, 1.5 mmol) were used to prepare 340 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 209 (14), 208 (100), 206 (6), 121 (11),
10 114 (5), 107 (5), 91 (6), 77 (12), 71 (8), 70 (16).

EXAMPLE 54: Preparation of N-(2-Hydroxy-2-methyl-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 101

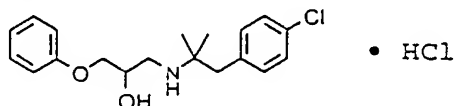


15 To a solution of 3-chloroperoxybenzoic acid (70% pure, 4.2 g, 17 mmol) in 40 mL of chloroform was added methallyl phenyl ether (2.5 g, 16.87 mmol). The mixture was stirred at room temperature for 5 hours then worked up by pouring into ether and sodium bicarbonate. The organic phase was washed
20 with sodium bisulfite, sodium bicarbonate, and sodium chloride, dried over anhydrous sodium sulfate and evaporated to give 2.3 g of 1,2-epoxy-2-methyl-3-phenoxypropane.

Using the method of Example 6, *supra*, 1,2-epoxy-2-methyl-3-phenoxypropane (0.092 g, 0.56 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.10 g, 0.56 mmol)
25 were used to prepare 120 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 328 (M-15,1), 223 (15), 222

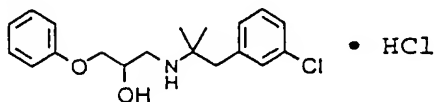
(100), 163 (13), 147 (13), 121 (21), 107 (11), 91 (9), 77 (13) 71 (12), 70 (49).

EXAMPLE 55: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-chlorophenyl)ethylamine Hydrochloride, Compound 103



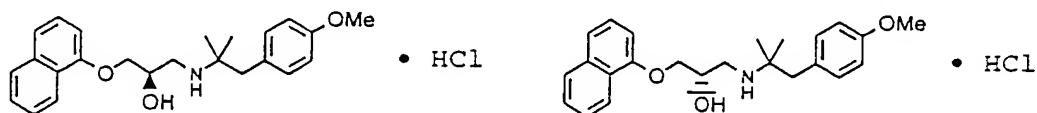
Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (950 mg, 6.3 mmol) and 1,1-dimethyl-2-(4-chlorophenyl)ethylamine (1.45 g, 7.9 mmol) were used to prepare 150 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 318 (M-15, 7), 209 (47), 208 (100), 127 (11), 125 (33), 114 (13), 107 (12), 77 (23), 71 (16), 70 (29), 58 (13).

EXAMPLE 56: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(3-chlorophenyl)ethylamine Hydrochloride, Compound 104



Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (1.3 g, 8.8 mmol) and 1,1-dimethyl-2-(4-chlorophenyl)ethylamine (2.0 g, 11 mmol) were used to prepare 338 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.) 318 (M-15, 1), 209 (14), 208 (100), 133 (4), 125 (12), 114 (6), 107 (6), 77 (10), 71 (8), 70 (15), 58 (5).

EXAMPLE 57: Resolution of the Enantiomers of (R)- and (S)- N-[2-Hydroxy-3-(1-naphthoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compounds 105 and 106

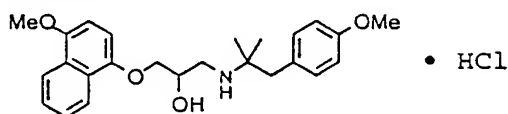


5 Compound 105

Compound 106

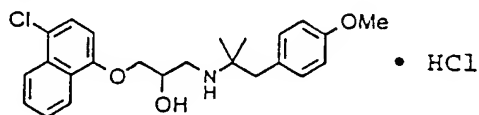
The free base (1.5 g) of compound 19 was chromatographed through ChiralCel OD (20 x 2.5 cm) using ethanol-hexane (1:4, plus 0.1% diethylamine) at 10 mL/min (270 nm). Chromatography of each enantiomer through Vydac C-18 (5 x 25
10 cm) using a gradient of 0.1% HCl to acetonitrile (50 mL/min., 264 nm) afforded the hydrochloride salt of compound 105 (464 mg) $[\alpha]_D^{26} = 15.3^\circ$ (c = 0.928, CHCl₃), m.p. 113-115°C and compound 106 (463 mg) $[\alpha]_D^{26} = -13.8^\circ$ (c = 0.926, CHCl₃).

EXAMPLE 58: Preparation of N-[2-Hydroxy-3-(4-methoxy-(1-naphthoxy))propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 107



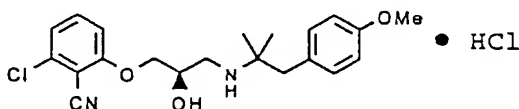
Using the method of Example 5, *supra*, 1,2-epoxy-3-[4-methoxy-(1-naphthoxy)]propane (462 mg, 2 mmol) and 1,1-
20 dimethyl-2-(4-methoxyphenyl)ethylamine (120 mg, 0.67 mmol) yielded, after preparative TLC and RP-HPLC, 116 mg of the title compound as a white solid: GC/EI-MS, m/z (rel. int.) 394 (M-15, 2), 288 (100), 731 (11), 121 (15), 71 (22).

EXAMPLE 59: Preparation of N-[2-Hydroxy-3-(4-chloro-(1-naphthoxy))propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 108



5 Using the method of Example 5, *supra*, 1,2-epoxy-3-[4-chloro-(1-naphthoxy)]propane (469 mg, 2 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (120 mg, 0.67 mmol) yielded, after preparative TLC and RP-HPLC, 131 mg of the title compound as a white solid: GC/EI-MS, *m/z* (rel. int.)
 10 414 (*M*⁺, 0.5), 398 (1), 292 (100), 121 (33), 71 (43).

EXAMPLE 60: Preparation of (R)-N-[2-Hydroxy-3-(3-chloro-2-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 109



15 To a solution of 18-crown-6 (3.96 g, 15 mmol) in 30 mL of acetonitrile were added dry potassium acetate (1.47 g, 15 mmol) and 2-chloro-6-fluorobenzonitrile (1.56 g, 10 mmol). The reaction was refluxed under nitrogen for 25 hours, then cooled to room temperature. Sodium hydroxide (2 mL of a 10 M
 20 solution, 20 mmol) and water (5 mL) were added, and the reaction stirred at room temperature for two hours. The acetonitrile was removed on a rotary evaporator, and the residue was taken up in ether and water. The basic aqueous layer was washed three times with ether. The aqueous layer
 25 was then made acidic with HCl, and the product extracted into ether. The ether layer was dried over anhydrous sodium

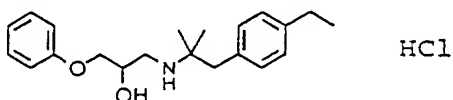
sulfate, filtered, and concentrated. The resulting solid was crystallized from water/methanol to yield 1.11 g of 3-chloro-2-cyanophenol.

3-Chloro-2-cyanophenol (0.55 g, 3.58 mmol) was dissolved in 10 mL of dimethylformamide, and the solution cooled to 0 °C. Sodium hydride (0.158 g, 3.94 mmol 60% in oil), washed with hexane and dimethylformamide, was added to cooled solution over a period of one minute. After stirring for 10 minutes at room temperature, (2R)-(-)-glycidyl 3-nitrobenzenesulfonate was added and stirred 16 hours. The reaction was poured into ether and dilute sodium hydroxide. The ether layer was separated and the aqueous layer extracted once more with ether. The combined ether extracts were washed with water and saturated brine, dried over anhydrous sodium sulfate, filtered, and concentrated to give 0.7 g of (R)-3-chloro-2-cyanophenyl glycidyl ether.

Using the method of Example 6, *supra*, (R)-3-chloro-2-cyanophenyl glycidyl ether (0.7 g, 3.34 mmol) and 1,1-dimethyl-(4-methoxyphenyl)ethylamine (0.72 g, 4.0 mmol) were used to prepare 570 mg of the title compound as a white solid: ¹H-NMR (CDCl₃) 9.65 (1H, br s), 8.2 (1H, br s), 7.4 (1H, t), 7.15 (2H, d), 7.03 (1H, d), 6.95 (1H, d), 6.8 (2H, d), 4.8 (1H, m), 4.3 (2H, d), 3.75 (3H, s), 3.4 (2H, m), 3.13 (2H, dd), 1.44 (3H, s), 1.40 (3H, s); ¹³C-NMR 161.9, 159.4, 138.2, 135.1, 132.4, 126.8, 122.8, 114.4, 114.2, 111.6, 104.0, 71.8, 66.0, 61.9, 55.8, 45.4, 43.9, 23.5, 23.3.

EXAMPLE 61: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-ethylphenyl)ethylamine Hydrochloride, Compound 110

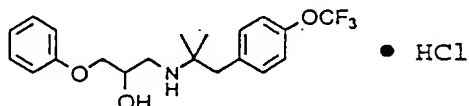
30



Using the method of Example 52, *supra*, 4-ethylbenzylamine (4.0 g, 29.6 mmol) was used to prepare 3.6 g of 1,1-dimethyl-2-(4-ethylphenyl)ethylamine. Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.43 g, 2.9 mmol) and 1,1-dimethyl-2-(4-ethylphenyl)ethylamine (0.5 g, 2.8 mmol) were used to prepare 600 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 313 (M-15, .1), 209 (23), 208 (100), 133 (5), 119 (12), 114 (6), 107 (5), 104 (7), 91 (6), 77 (10), 71 (8), 70 (12).

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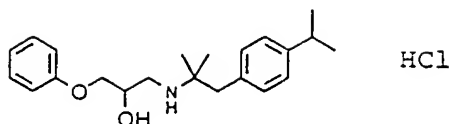
EXAMPLE 62: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-trifluoromethoxyphenyl)ethylamine Hydrochloride, Compound 111



15 Using the method of Example 52, *supra*, 4-trifluoromethoxybenzylamine (2.0 g, 10.5 mmol) was used to prepare 2.2 g of 1,1-dimethyl-2-(4-trifluoromethoxyphenyl)ethylamine.

Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.12 g, 0.8 mmol) and 1,1-dimethyl-2-(4-trifluoromethoxyphenyl)ethylamine (0.175 g, 0.8 mmol) were used to prepare 15 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 368 (M-15,2), 209 (39), 208 (100), 175 (20), 133 (5), 114 (5), 107 (6), 77 (11), 71 (7), 70 (12), 58 (5).

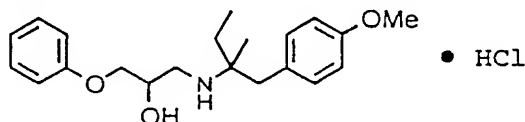
25 EXAMPLE 63: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-isopropylphenyl)ethylamine Hydrochloride, Compound 112



Using the method of Example 52, *supra*,
 4-isopropylbenzylamine (4.89 g, 32.8 mmol) was used to
 prepare 4.1 g of 1,1-dimethyl-2-(4-isopropylphenyl)-
 5 ethylamine. Using the method of Example 6, *supra*,
 1,2-epoxy-3-phenoxypropane (0.173 g, 1.15 mmol) and
 1,1-dimethyl-2-(4-isopropylphenyl)ethylamine (0.275 g, 1.44
 mmol) were used to prepare 89 mg of the title compound as a
 white solid: GC/EI-MS, *m/z*, (rel. int.) 326 (M-15,1), 209
 10 (14), 208 (100), 133 (9), 117 (5), 114 (5), 105 (5), 91 (6),
 77 (8), 71 (8), 70 (13), 58 (5).

EXAMPLE 64: Preparation of N-(2-Hydroxy-3-
phenoxypropyl)-1-ethyl-1-methyl-2-(4-methoxyphenyl)ethylamine
Hydrochloride, Compound 113

15



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4-Hydroxybenzyl alcohol (0.35 g, 2.82 mmol) and
 tetrabutylammonium fluoride (0.147 g, 0.56 mmol) were
 dissolved in 3 mL of 2-nitrobutane and heated to 130-145 °C
 under nitrogen for 20 hours. The reaction mixture was cooled
 and partitioned between water and ether. The ether layer was
 separated, washed with saturated brine, dried over anhydrous
 sodium sulfate, and concentrated. The crude material was
 purified by preparative TLC using ethyl acetate/hexane as the
 elutant. The yield of 1-ethyl-1-methyl-2-(4-hydroxyphenyl)-
 25 nitroethane was 0.21 grams.

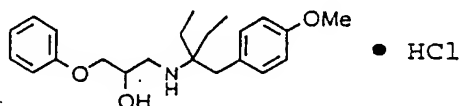
To a suspension of 40% (wt/wt) potassium fluoride on alumina (0.73 g, 5 mmol) in 3 mL of acetonitrile were added 1-ethyl-1-methyl-2-(4-hydroxyphenyl)nitroethane (0.21 g, 1.0 mmol) and iodomethane (0.21 g, 1.5 mmol). The reaction was stirred at room temperature for 4 days and then filtered and rinsed with acetonitrile. The acetonitrile was removed on a rotary evaporator, and the residue was partitioned between ether and water. The ether layer was separated, washed with sodium bisulfite, sodium carbonate, and saturated brine, then dried over anhydrous sodium sulfate and concentrated. The yield of 1-ethyl-1-methyl-2-(4-methoxyphenyl)nitroethane was 0.183 g.

Nickel chloride monohydrate (0.107 g, 0.404 mmol) was dissolved in 5 mL of methanol, followed by the addition of sodium borohydride (0.05 g, 1.2 mmol). After stirring for 5 minutes, 1-ethyl-1-methyl-2-(4-methoxyphenyl)nitroethane (0.18 g, 0.807 mmol) in 3 mL of methanol was added, and stirred for 5 minutes. Sodium borohydride (0.11 g, 2.83 mmol) was then added in portions over 5 minutes. The reaction was then stirred overnight under a hydrogen balloon. The reaction mixture was filtered, and the methanol was removed on a rotary evaporator. The residue was taken up in ether and dilute sodium hydroxide. The ether layer was separated, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The yield of 1-ethyl-1-methyl-2-(4-methoxy-phenyl)ethylamine was 0.127 grams.

Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.086 g, 0.57 mmol) and 1-ethyl-1-methyl-2-(4-methoxyphenyl)ethylamine (0.11 g, 0.57 mmol) were used to prepare 90 mg of the title compound as a white solid:
GC/EI-MS, *m/z*, (rel. int.) 314 (M-29,2), 223 (15), 222 (100),

128 (6), 121 (20), 107 (5), 84 (10), 78 (5), 77 (12), 72 (5),
56 (7).

EXAMPLE 65: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-diethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 114



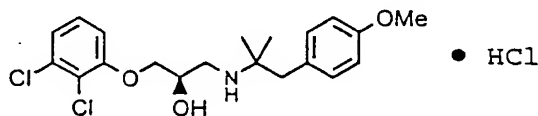
Anhydrous cerium (III) chloride (13.6 g, 55.2 mmol) was suspended in 80 mL of dry tetrahydrofuran, and stirred under nitrogen for 16 hours. This suspension was cooled in an ice bath, and ethylmagnesium chloride (27.6 mL, 55.18 mmol, 2 M solution in tetrahydrofuran) was added over 5 minutes. After stirring for 1 hour, methyl 4-methoxyphenylacetate (3.98 g, 22.07 mmol) was added to the suspension and stirred for another 2 hours. The reaction was then partitioned between ether and saturated ammonium chloride. The ether layer was separated, washed with dilute HCl, water, and saturated brine, dried over anhydrous sodium sulfate, and concentrated. The yield of 1,1-diethyl-2-(4-methoxyphenyl)ethanol was 4.65 g.

20 Powdered sodium cyanide (1.18 g, 24 mmol) was placed in
a flask and covered with 5.5 mL of acetic acid. A mixture of
sulfuric acid (3 mL) and acetic acid (2.75 mL) was cooled to
0 °C and then added to the cyanide suspension over a period
of 3 minutes. The mixture was stirred for 30 minutes at room
25 temperature, followed by the addition of 1,1-diethyl-2-(4-
methoxyphenyl)ethanol (4.6 g, 22 mmol). The mixture was
stirred overnight then poured into ice and sodium hydroxide.
The product was extracted with ether, and the ether layer
dried over anhydrous sodium sulfate, and concentrated. The

residue was suspended in 20% sodium hydroxide and refluxed overnight under nitrogen. The reaction was cooled, diluted with water, and extracted with ether. The ether layer was separated, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by reversed-phase HPLC (C-18 using 0.1% HCl/acetonitrile as the elutant) to give 1.91 g of 1,1-diethyl-2-(4-methoxyphenyl)ethylamine.

Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.15 g, 1.0 mmol) and 1,1-diethyl-2-(4-methoxyphenyl)ethylamine (0.249 g, 1.2 mmol) were used to prepare 244 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 328 (M-29,6), 237 (17), 236 (100), 121 (22), 106 (5), 98 (7), 78 (5), 77 (11), 70 (7), 56 (5).

EXAMPLE 66: Preparation of (R)-N-[2-Hydroxy-3-(2,3-dichlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 115

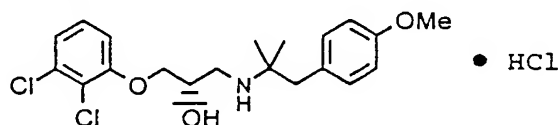


2,3-Dichlorophenol (0.69 g, 4.24 mmol) was dissolved in 15 mL of acetone, followed by the addition of powdered potassium carbonate (1.6 g, 11.57 mmol). This mixture was stirred for 2 hours then (2R)-(-)-glycidyl 3-nitrobenzenesulfonate was added and stirred overnight. The reaction was worked up by pouring into water and ether. The ether layer was separated, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The yield of (R)-2,3-dichlorophenyl glycidyl ether was 0.837 g.

Using the method of Example 6, *supra*, (R)-2,3-dichlorophenyl glycidyl ether (0.837 g, 3.82 mmol) and

1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.75 g, 4.18 mmol) were used to prepare 860 mg of the title compound as a white solid: GC/EI-MS, m/z , (rel. int.) 382 (M-15, .1), 280 (11), 278 (64), 277 (16), 276 (100), 163 (10), 121 (35), 77 (10), 71 (24), 70 (27), 58 (12).

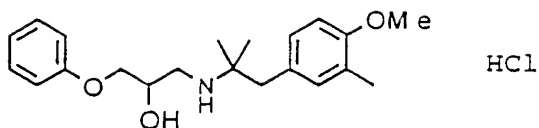
EXAMPLE 67: Preparation of (S)-N-[2-Hydroxy-3-(2,3-dichlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 116



2,3-Dichlorophenol (0.69 g, 4.24 mmol) was dissolved in 15 mL of acetone, followed by the addition of powdered potassium carbonate (1.6 g, 11.57 mmol). This mixture was stirred for 2 hours then (2S)-(+)-glycidyl 3-nitrobenzenesulfonate was added and stirred overnight. The reaction was worked up by pouring into water and ether. The ether layer was separated, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The yield of (S)-2,3-dichlorophenyl glycidyl ether was 0.84 g.

Using the method of Example 6, *supra*, (S)-2,3-dichlorophenyl glycidyl ether (0.84 g, 3.82 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.75 g, 4.18 mmol) were used to prepare 860 mg of the title compound as a white solid: GC/EI-MS, m/z , (rel. int.) 382 (M-15, .1), 280 (10), 279 (9), 278 (63), 276 (100), 163 (11), 121 (28), 77 (7), 71 (21), 70 (23), 58 (10).

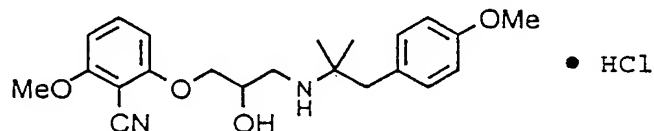
EXAMPLE 68: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-methoxy-3-methylphenyl)ethylamine Hydrochloride, Compound 117



Using the method of Example 64, *supra*, 4-hydroxy-3-methylbenzyl alcohol (1.0 g, 7.25 mmol), 2-nitropropane (5 mL), and tetrabutylammonium fluoride (0.38 g, 0.145 mmol) were used to prepare 0.8 g of 1,1-dimethyl-2-(4-methoxy-3-methylphenyl)ethylamine.

Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.151 g, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxy-3-methylphenyl)ethylamine (0.2 g, 1.0 mmol) were used to prepare 130 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 328 (M-15, .1), 209 (14), 208 (100), 177 (5), 135 (14), 114 (5), 91 (6), 76 (9), 71 (8), 70 (13), 58 (5).

EXAMPLE 69: Preparation of N-[2-Hydroxy-3-(2-cyano-3-methoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 118



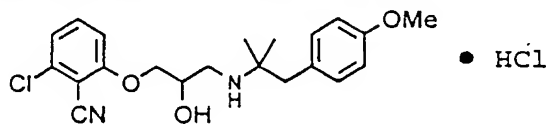
Powdered sodium cyanide (9.0 g, 184 mmol) and 2,6-dimethoxybenzonitrile were added to 50 mL of dimethylsulfoxide and heated to 145 °C for 110 min under nitrogen. The reaction was cooled and poured into ether and dilute HCl. The ether layer was separated, washed twice with dilute acid, once with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The yield of 2-cyano-3-methoxyphenol was 8.1 g.

2-Cyano-3-methoxyphenol (1 g, 6.7 mmol) and powdered potassium carbonate (2.78 g, 20.1 mmol) were stirred in 15 mL of acetone for 5 minutes, followed by addition of epibromohydrin (1.38 g, 10.1 mmol). The mixture was stirred for 72 hours then poured into water/ether. The ether layer was separated, washed with sodium carbonate and saturated brine, dried over anhydrous sodium sulfate, and concentrated. The resulting crude solid was triturated with ether/hexane, filtered, and dried under vacuum to give 0.44 g of 2-cyano-3-methoxyphenyl glycidyl ether.

Using the method of Example 6, *supra*, 2-cyano-3-methoxyphenyl glycidyl ether (0.205 g, 1.0 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.215 g, 1.2 mmol) were used to prepare 265 mg of the title compound as a white solid: ¹H-NMR (CDCl₃) δ 9.6 (1H, br s), 8.2 (1H, br s), 7.4 (1H, t), 7.15 (2H, d), 6.8 (2H, d), 6.6 (1H, d), 6.53 (1H, d), 4.75 (1H, m), 4.25 (2H, m), 3.87 (3H, s), 3.77 (3H, s), 3.43 (2H, m), 3.12 (2H, dd), 1.45 (3H, s), 1.41 (3H, s). ¹³C-NMR δ 162.9, 162, 159.3, 135.5, 132.4, 127, 114.7, 114.4, 105.6, 104.6, 92.2, 71.4, 66.1, 61.9, 56.8, 55.8, 45.4, 43.8, 23.5, 23.3.

EXAMPLE 70: Preparation of N-[2-Hydroxy-3-(3-chloro-2-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 119

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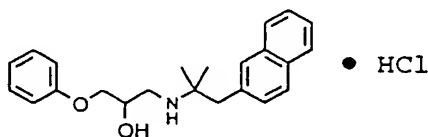


Using the method of Example 69, *supra*, 3-chloro-2-cyanophenol (made using the method of Example 60, *supra*) (0.48 g, 3.13 mmol), potassium carbonate (1.3 g, 9.38 mmol),

and epibromohydrin (0.86 g, 6.25 mmol) were used to prepare 93 mg of 3-chloro-2-cyanophenyl glycidyl ether.

Using the method of Example 6, *supra*, 3-chloro-2-cyanophenyl glycidyl ether (0.093 g, 0.44 mmol) and 1,1-dimethyl-(4-methoxyphenyl)ethylamine (0.095 g, 0.53 mmol) were used to prepare 134 mg of the title compound as a white solid: ¹H-NMR (CDCl₃) δ 9.68 (1H, br s), 8.2 (1H, br s), 7.4 (1H, t), 7.15 (2H, d), 7.03 (1H, d), 6.95 (1H, d), 6.8 (2H, d), 5.7 (1H, br s), 4.8 (1H, m), 4.3 (2H, d), 3.75 (3H, s), 3.4 (2H, m), 3.13 (2H, dd), 1.44 (3H, s), 1.40 (3H, s).

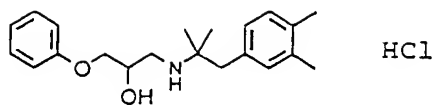
EXAMPLE 71: Preparation of N-(2-Hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(2-naphthyl)ethylamine Hydrochloride, Compound 120



Using the method of Example 52, *supra*, 2-amino-methylnaphthalene (2.51 g, 16 mmol) was used to prepare 1.9 g of 1,1-dimethyl-2-(2-naphthyl)ethylamine.

Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.163 g, 1.1 mmol) and 1,1-dimethyl-2-(2-naphthyl)ethylamine (0.26 g, 1.3 mmol) were used to prepare 243 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 334 (M-15, .1), 209, (14), 208 (100), 141 (16), 115 (7), 76 (5), 70 (7).

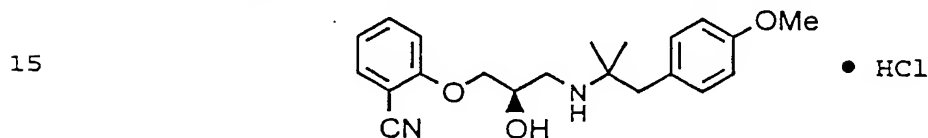
EXAMPLE 72: Preparation of N-(2-Hydroxy-3-phenoxy)propyl)-1,1-dimethyl-2-(3,4-dimethylphenyl)ethylamine Hydrochloride, Compound 121



Using the method of Example 52, *supra*, 3,4-dimethylbenzylamine (5 g, 37 mmol) was used to prepare 2.29 g of 1,1-dimethyl-2-(3,4-dimethylphenyl)ethylamine.

5 Using the method of Example 6, *supra*, 1,2-epoxy-3-phenoxypropane (0.165 g, 1.1 mmol) and 1,1-dimethyl-2-(3,4-dimethylphenyl)ethylamine (0.22 g, 1.2 mmol) were used to prepare 268 mg of the title compound as a white solid: GC/EI-MS, *m/z*, (rel. int.) 312 (M-15,1), 209 (14), 208 (100),
 10 133 (5), 119 (13), 114 (5), 107 (5), 91 (5), 76 (10), 71 (8), 70 (14), 58 (6).

EXAMPLE 73: Preparation of (R)-N-[2-Hydroxy-3-(2-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 122



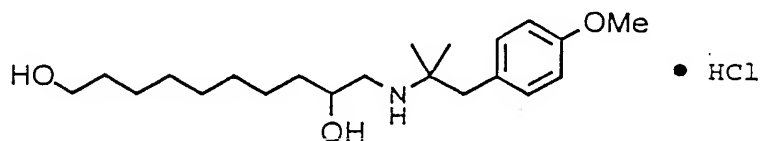
15 Using the method of Example 60, *supra*, 2-cyanophenol (0.54 g, 4.5 mmol), sodium hydride (0.188 g, 4.7 mmol), and (2R)-(-)-glycidyl 3-nitrobenzenesulfonate (1.06 g, 4.1 mmol) were used to prepare 350 mg of (R)-2-cyanophenyl glycidyl
 20 ether.

Using the method of Example 6, *supra*, (R)-2-cyanophenyl glycidyl ether (0.35 g, 2.0 mmol) and 1,1-dimethyl-(4-methoxyphenyl)ethylamine (0.35 g, 1.96 mmol) were used to prepare 600 mg of the title compound as a white solid: ¹H-NMR
 25 (CDCl₃) δ 9.7 (1H, br s), 8.2 (1H, br s), 7.5 (2H, m), 7.15 (2H, d), 7.0 (2H, m), 6.8 (2H, d), 4.8 (1H, br m), 4.25 (2H,

m), 3.75 (3H, s), 3.45 (2H, m), 3.12 (2H, dd), 1.45 (3H, s), 1.41 (3H, s).

EXAMPLE 74: Preparation of N-(2,10-Dihydroxydecyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride.

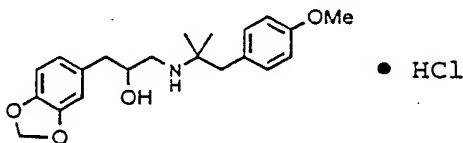
5 Compound 123



Using the method of Example 9, *supra*, 1,2-epoxy-10-hydroxydecane (172 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (230 mg, 1.5 mmol) were used to
10 prepare the hydrochloride salt of the title compound. MPLC of the free amine (silica gel, 1% MeOH/CHCl₃), followed by treatment with an excess of 1 M HCl/ether, yielded 130 mg of the title compound as a white powder: GC/EI-MS, *m/z* (rel. int.) 336 (*M*⁺ - 15, .1), 231 (14), 230 (100), 212 (9), 163
15 (10), 122 (5), 121 (42), 91 (8), 78 (6), 77 (5), 71 (13), 70 (11), 58 (8), 55 (8), 41 (6).

EXAMPLE 75: Preparation of N-[2-hydroxy-3-(3,4-methylene-dioxyphenyl)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride. Compound 124

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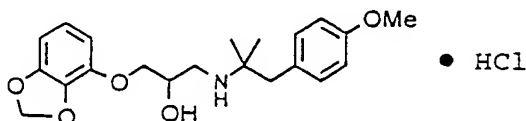


Safrole oxide was prepared by stirring a solution of safrole (1.48 mL, 10 mmol) and *m*-chloroperoxybenzoic acid (2.70 g, 11 mmol) in methylene dichloride (25 mL) overnight. The reaction was quenched by pouring into water (50 mL). The

aqueous was extracted with ether (3 x 25 mL). The organic layers were combined and washed with 10% aqueous sodium sulfate (2 x 25 mL), saturated aqueous sodium bicarbonate (3 x 25 mL), and brine (25 mL). The organic phase was dried
 5 over magnesium sulfate and the solvents were removed in vacuo. The resulting yellow oil was used without further purification.

To a solution of safrole oxide (196 mg, 1.1 mmol) in acetonitrile (1.0 mL) was added lithium perchlorate (107 mg).
 10 The solution was stirred at ambient temperature to dissolve all of the solid lithium perchlorate. To this solution was added 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (180 mg, 1.0 mmol) and the reaction was stirred at 50 °C overnight. To the cooled reaction mixture was added water (5 mL) and was
 15 subsequently extracted with methylene dichloride (3 x 1 mL). The combined organic phases were washed with water (1 mL) and brine (1 mL) and dried over magnesium sulfate. The crude orange oil was purified (MPLC, silica gel, 1% MeOH/CHCl₃) and dissolved in methylene dichloride (5 mL). The hydrochloride
 20 salt was prepared by adding an excess of 1 M HCl/ether. The solvents were removed in vacuo to yield 139 mg of thick oil: GC/EI-MS, *m/z* (rel. int.) 342 (M⁺, .1), 237 (15), 236 (100), 163 (5), 136 (6), 135 (61), 121 (23), 78 (7), 77 (13), 70 (15), 58 (7).

25 EXAMPLE 76: Preparation of N-[2-hydroxy-3-(3,4-methylene-dioxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)-ethylamine Hydrochloride, Compound 125

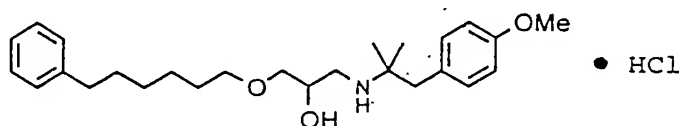


Using the method of Example 9, *supra*, 1,2-epoxy-3-(3,4-methylenedioxyphenoxy)propane (194 mg, 1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (180 mg, 1 mmol) were used to prepare the hydrochloride salt of the title compound.

5 MPLC of the free amine (silica gel, 1% MeOH/CHCl₃), followed by treatment with an excess of 1 M HCl/ether, yielded 88 mg of a white powder: GC/EI-MS, *m/z* (rel. int.) 374 (M⁺, .0), 253 (15), 252 (100), 137 (8), 121 (16), 114 (8), 71 (12), 70 (8).

EXAMPLE 77: Preparation of N-[2-hydroxy-3-(6-phenyl-hexanoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

10 Compound 126

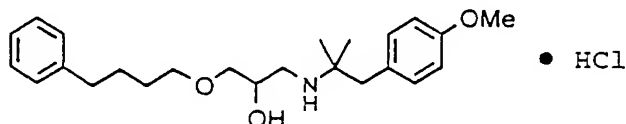


Using the method of Example 9, *supra*, 6-phenyl-hexylglycidyl ether (337 mg, 1.44 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (180 mg, 1 mmol) were used to prepare the title compound. Preparative TLC (20 cm x 20 cm x 2 mm silica, eluted with 1% MeOH/CHCl₃) was used to purify the material and yielded 275 mg of free base: GC/EI-MS, *m/z* (rel. int.) 398 (M⁺ - 15, .1), 293 (21), 292 (100), 163 (10), 121 (19), 114 (9), 91 (5), 90 (45), 71 (13), 70 (14), 58 (9).

20

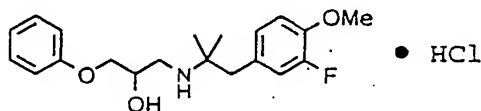
EXAMPLE 78: Preparation of N-[2-hydroxy-3-(4-phenyl-butanoxyl)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 127



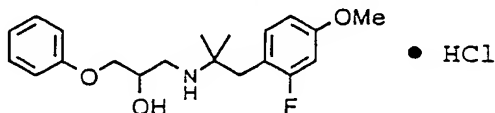
Using the method of Example 9, *supra*, 4-phenylbutyl glycidyl ether (348 mg, 1.5 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (268 mg, 1.5 mmol) were used to prepare the title compound. Preparative TLC (20 cm x 20 cm x 2 mm silica, eluted with 5% MeOH/CHCl₃) was used to purify the material and yielded 275 mg of free base: GC/EI-MS, *m/z* (rel. int.) 370 (*M*⁺ - 15, .1), 265 (19), 264 (100), 163 (11), 121 (19), 114 (9), 90 (43), 71 (10), 70 (12), 58 (7).

EXAMPLE 79: Preparation of N-(2-hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(3-fluoro-4-methoxyphenyl)ethylamine, Compound 128



Using the method of Example 6, *supra*, phenyl glycidyl ether (150 mg, 1.1 mmol) and 1,1-dimethyl-2-(3-fluoro-4-methoxyphenyl)ethylamine (197 mg, 1 mmol) were used to prepare the title compound. The title compound crystallized on standing to yield 169 mg of small crystals: GC/EI-MS, *m/z* (rel. int.) 332 (*M*⁺ - 15, .1), 209 (16), 208 (100), 139 (13), 133 (5), 114 (7), 107 (6), 77 (11), 71 (12), 70 (20), 58 (9).

EXAMPLE 80: Preparation of N-(2-hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(2-fluoro-4-methoxyphenyl)ethylamine Hydrochloride, Compound 129

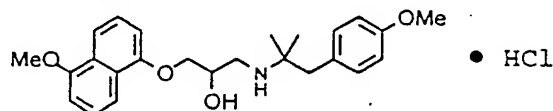


Using the method of Example 6, *supra*, phenyl glycidyl ether (150 mg, 1.1 mmol) and 1,1-dimethyl-2-(2-fluoro-

4-methoxyphenyl)ethylamine (197 mg, 1 mmol) were used to prepare the title compound. Preparative HPLC (C_{18} reversed-phase, eluted with 1% HCl/ CH_3CN gradient) was used to purify the compound, yielding 301 mg of a white powder:

- 5 GC/EI-MS, m/z (rel. int.) 332 ($M^+ - 15, 1$), 209 (15), 208 (100), 139 (14), 114 (5), 107 (5), 77 (6), 71 (7), 70 (13), 58 (5).

EXAMPLE 81: Preparation of N-[2-hydroxy-3-(5-methoxy-1-naphthoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
 10 Hydrochloride, Compound 130



- Potassium carbonate (13 mmol) was added to a solution of 1,5-dihydroxy naphthalene (6.2 mmol) in acetone in a sealed vacuum tube. The tube was heated to 70 °C for 30 minutes.
 15 Iodomethane (9.4 mmol) was added and the tube heated to 70 °C overnight.

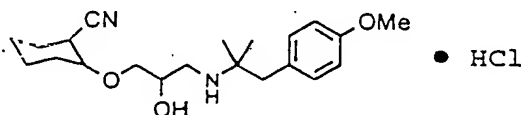
- The reaction mixture was partitioned between ether and 10% aqueous HCl. The ether layer was separated and extracted with 0.5 M KOH. The water layer was separated and acidified
 20 with 10% aqueous HCl and extracted into ether. The ether layer was separated and dried over magnesium sulfate and evaporated to a solid. The solid was purified using reverse phase HPLC using a acetonitrile/0.1% HCl gradient yielding 179 mg 1-hydroxy-5-methoxy naphthalene.

- 25 Sodium hydride (60% suspension in mineral oil, 1 mmol) was added to a solution of 1-hydroxy-5-methoxy naphthalene (1 mmol) and stirred 10 minutes. Epichlorohydrin (1 mmol) was added and the reaction stirred at 70 °C for 72 hours. The reaction mixture was diluted with 1 liter of saturated sodium

chloride solution and extracted into ether. The ether layer was then washed with water, separated and dried over anhydrous sodium sulfate and evaporated to give 5-methoxy-napthalene glycidyl ether.

- 5 Using the method of Example 6, *supra*, 5-methoxy-napthalene glycidyl ether (1 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1 mmol) yielded 161 mg of the title compound as a white solid: GC /EI-MS, m/z (rel. int.) 360 (M+, 1), 289 (18), 288 (100), 173 (8), 121 (20), 71 (18),
10 70 (12).

EXAMPLE 82: Preparation of N-[2-hydroxy-3-(2-cyano-cyclohexyloxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride, Compound 131

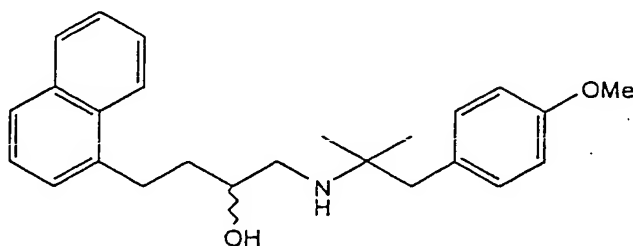


- 15 Sodium hydride (60% suspension in mineral oil, 60 mg, 1.59 mmol) was added to a solution of trans-2-cyano-cyclohexanol in N,N-dimethylformamide (2.0 mL) and stirred for 10 minutes at room temperature. Epibromohydrin (0.22 g, 1.59 mmol) was then added and the reaction stirred for an
20 additional 3 hours. The solution was partitioned between diethyl ether/water, and the layers separated. The ether layer was washed with water (3 X 100 mL) and dried over magnesium sulfate and evaporated to give 0.17 g of 2-cyanocyclohexyl glycidyl ether.
- 25 Using the method of Example 6, *supra*, 2-cyanocyclohexyl glycidyl ether (0.94 mmol) and 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (1.17 mmol) yielded 55 mg of the title compound as a white solid: GC /EI-MS, m/z (rel. int.) 360

(M+, 1), 239 (100), 121 (22), 240 (14), 70 (11), 163 (8), 71 (8), 81 (7).

EXAMPLE 83: Synthesis of (R/S)-1-[[2,2-dimethyl-(4-methoxy) phenethyl]amino-2-hydroxy-4(1'-naphthyl)-butane.

5 Compound 162

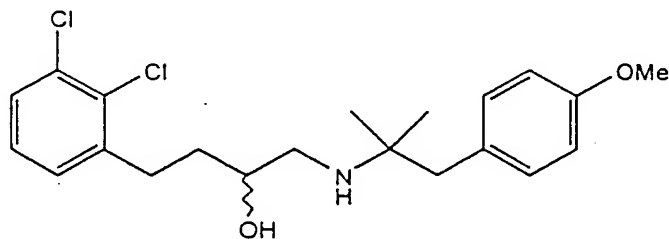


A solution of 1-chloromethylnaphthalene (750 uL, 5 mmol, Aldrich) in anhydrous ether (10 mL) was added dropwise to 25 mL of allyl magnesium bromide (1 M in ether) over 30 minutes. 10 The resulting mixture was heated at reflux for 14 hours. After cooling the reaction to room temperature, it was quenched with 25 mL of saturated NH_4Cl (aqueous). The layers were separated and the organic layer was washed with brine, dried over Na_2SO_4 , filtered and evaporated to give 1 g of 1- 15 but-3-enyl-naphthalene that was carried without further purification.

1-But-3-enyl-naphthalene (1 g) from above was added to a solution of 50% mCPBA (2.1 g) in CH_2Cl_2 (50 mL) and the reaction stirred at room temperature for 48 hours. The 20 material was diluted with CH_2Cl_2 , and was extracted with sodium sulfite (aqueous) and NaHCO_3 (aqueous), dried over MgSO_4 , filtered and evaporated to give 1-[(2-oxoaryl)ethyl]-naphthalene (1 g) that was carried without further purification.

A solution of 1-[(2-oxoaryl)ethyl]-naphthalene (1 g) and 1,1-dimethyl-2-(4-methoxyphenyl) ethylamine (985 mg, 5.5 mmol) in ethanol (25 mL) was heated at reflux for 12 hours: The reaction was evaporated and the residue dissolved in 4 N HCl/dioxane. Upon addition of ether, crystals formed and were subsequently collected and dried in a vacuum oven to give 1.4 g of (R/S)-1-[[2,2-dimethyl-(4'methoxy)-phenethyl]]amino-2-hydroxy-4(1'-naphthyl)-butane. ESMS [(M+H)⁺= 378, ¹H NMR (CDCl₃, 360MHz) @300°K δ 8.06 (1H, d of d), 7.83 (1H, d of d), 7.78-7.61 (2H, m), 7.49-7.35 (3H, m), 7.09-7.02 (2H, m), 6.84-6.79 (2H, m), 3.76 (3H, s), 3.61 (1H, m), 3.33-3.09 (2H, m), 2.77-2.72 (1H, d of d), 2.62 (2H, s), 2.47-2.42 (1H, m), 1.85-1.82 (2H, m), 1.04 (6H, s).

Example 84 Synthesis of (R/S)-1-[[2,2-dimethyl-(4'methoxy)phenethyl]]amino-2-hydroxy-4(1'-(2,3-dichlorophenyl))-butane, Compound 163

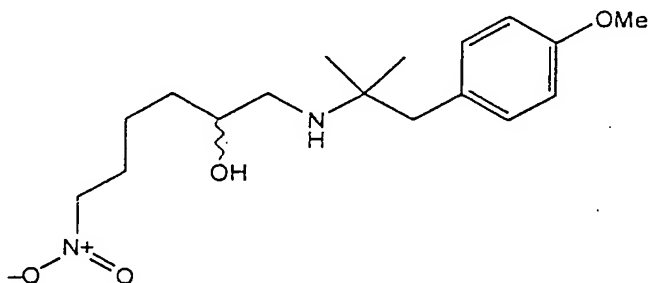


Starting from 2,3-dichlorobenzylchloride (1g, 5 mmol) and following the three step procedure described in Example 83, 660 mg of (R/S)-1-[[2,2-dimethyl-(4'methoxy)-phenethyl]]amino-2-hydroxy-4(1'-(2,3-dichlorophenyl))-butane was synthesized and isolated as white crystals. ESMS [M+H]⁺=396 ¹H NMR (CDCl₃, 360 MHz) @ 300°K δ 7.3 (1H, d of d), 7.18 (1H, d of d), 7.10-7.03 (3H, m), 6.82-6.80 (2H, m), 3.78

(3H, s), 3.47 (1H, m), 2.97-2.83 (2H, m), 2.74 (1H, m) 2.60 (2H, s), 2.43-2.37 (1H, m), 1.71 (2H, m), 1.04 (6H, s).

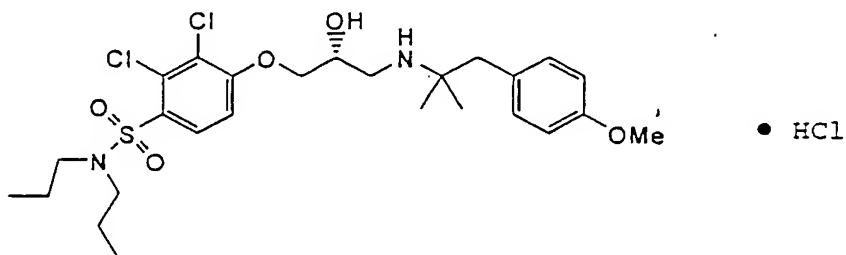
Example 85: Synthesis of (R/S)-1-nitro-5-hydroxy-6-[1,1-dimethyl-2-(4-methoxyphenyl)]hexane, Compound 164

5



Starting from 6-nitro-1-hexene (1 g, 7.75 mmol) and following the two last steps described in Example 83, 1 g of (R/S)-1-nitro-5-hydroxy-6-[1,1-dimethyl-2(4-methoxyphenyl)]hexane was synthesized and isolated as tan crystals. ESMS [M+H]⁺=325 ¹H NMR (CDCl₃, 360 MHz) @ 300°K δ 7.00 (4H, d of d), 4.37 (2H, t), 3.77 (3H, s), 3.49 (1H, m), 2.74 (1H, d of d), 2.61 (2H, s), 2.35 (1H, m) 2.03 (2H, m), 1.60-1.43 (4H, m) 1.08 (6H, s).

Example 86: N-[2(R)-Hydroxy-3-[(2,3-dichloro-4-dipropylsulfamoyl)phenoxy]-1-propyl]-N-[1,1-dimethyl-2-(4-methoxyphenyl)ethyl]amine hydrochloride salt
Compound 165



a) (2,3-Dichloro-4-methoxy)phenylsufonylchloride
 2,3-dichloroanisole (Aldrich, 9.0 g, 50.8 mmol) was utilized
 in the method of H. Harada et al., *Chem Pharm Bull* (1987)
 35(8) 3195-3214 to give the title compound as a white solid
 5 (13.3, 95%).

b) N,N-Dipropyl-(2,3-dichloro-4-methoxy)-
 phenylsulfonamide.

The compound of Example 86a (8.0 g, 29.0 mmol) was
 dissolved in CH_2Cl_2 (200 mL) and dipropylamine (11.9 mL, 87.1
 10 mmol) in EtOH (40 mL) was added at -20°C . The ice bath was
 removed and the mixture stirred 1.5 hours. The mixture was
 poured into H_2O and extracted with CH_2Cl_2 . The combined
 organic extracts were washed with H_2O and brine, concentrated
 in vacuo and azeotroped with toluene to yield the title
 15 compound as a light brown-tinted oil (9.8 g, 100%). ^1H NMR
 (400 MHz, CDCl_3) δ 8.05 (d, $J=10$ Hz, 1H), 6.92 (d, $J=10$ Hz,
 1H), 4.00 (s, 3H), 3.23 (t, $J=8, 17$ Hz, 4H), 1.52 (m, 4H),
 0.83 (t, $J=8, 13$ Hz, 6H).

c) N,N-Dipropyl-2,3-dichloro-4-hydroxyphenylsulfonamide.

20 The compound from Example 86b (10.0 g, 29.4 mmol), I_2
 (14.9 g, 58.8 mmol) and trimethylphenylsilane (15.1 mL, 88.2
 mmol) were stirred together and heated to 110°C for 18 hours.
 The mixture was poured into aqueous $\text{Na}_2\text{S}_2\text{O}_3$, extracted with
 EtOAc, dried (MgSO_4), concentrated to dryness in vacuo and
 25 purified by column chromatography (silica gel, 40% EtOAc in
 hexanes) to give a clear oil (9.2 g, 86%). ^1H NMR (400 MHz,
 CDCl_3) δ 7.98 (d, $J=10$ Hz, 1H), 7.05 (d, $J=10\text{Hz}$, 1H), 6.29
 (brs, 1H), 3.24 (t, $J=9, 18$ Hz, 4H), 1.56 (m, 4H), 0.84 (t,
 $J=8, 14$ Hz, 6H).

30 d) [2,3-Dichloro-4-(N,N-dipropylsulfamoyl)]phenyl
 glycidyl ether.

The compound of Example 86c (5.0 g, 15.3 mmol), K_2CO_3
 (6.4 g, 46.0 mmol), and (2R)-(-)-glycidyl 3-nitrobenzene-

sulfonate (5.6 g, 15.3 mmol) were heated in acetone (250 mL) to reflux 18 hours. The solvent was concentrated in vacuo to half volume, poured into H₂O, extracted with EtOAc, the combined organic extracts were dried (MgSO₄), evaporated and
 5 purified by column chromatography (silica gel, 40% EtOAc in hexanes) to give the title compound as a clear oil (4.9 g, 84%). ¹H NMR (400 MHz, CDCl₃) δ 8.04 (d, J=8 Hz, 1H), 6.94 (d, J=8 Hz, 1H), 4.45 (dd, J=1, 9 Hz, 1H), 4.11 (dd, J=7, 11 Hz, 1H), 3.44 (m, 1H), 3.24 (t, J=9, 18 Hz, 4H), 2.97 (m, 1H),
 10 2.87 (m, 1H), 1.52 (m, 4H), 0.84 (t, J=6, 14 Hz, 6H)

e) N-[2(R)-Hydroxy-3-[(2,3-dichloro-4-dipropylsulfamoyl)phenoxy]-1-propyl]-N-[1,1-dimethyl-2-(4-methoxyphenyl)ethyl]amine hydrochloride salt.

The compound from Example 86d (1.6 g, 4.2 mmol),
 15 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine (0.75 g, 4.2 mmol) and LiClO₄ (0.89 g, 8.4 mmol) were dissolved in CH₃CN (150 mL) and refluxed 18 hours. The mixture was concentrated in vacuo and purified by column chromatography (silica gel, 8% MeOH in CH₂Cl₂) to yield the title compound as a white solid (1.0 g,
 20 44%). This was converted to the HCl salt by adding 1.7 mL of 1M HCl in MeOH, stirring 5 minutes, concentrating in vacuo, azeotroping with toluene then CH₂Cl₂. MS (ES) m/e 561.1 [M+H⁺]; ¹H NMR (400 MHz, CDCl₃) δ 9.94 (bs, 1H), 8.02 (d, J=8 Hz, 1H), 7.14 (d, J=8 Hz, 2H), 6.96 (d, J=8 Hz, 1H), 6.86
 25 (d, J=7 Hz, 1H), 4.78 (m, 1H), 4.30 (m, 1H), 4.20 (m, 1H), 3.80 (s, 3H), 3.68 (m, 2H), 3.43 (bs, 1H), 3.23 (t, J=7, 14 Hz, 4H), 3.12 (m, 2H), 1.50 (q, J=5, 13 Hz, 4H), 1.43 (s, 3H), 1.38 (s, 3H), 0.83 (t, J=8, 13, 6H).

EXAMPLE 87: Additional compounds

30 Additional compounds were synthesized using techniques along lines as those described above. Examples of such compounds include the following:

Compound 1: *N*-[2-hydroxy-3-(2-hydroxybenzimidazol-4-oxyl)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 14: (*R*)-*N*-[2-hydroxy-3-(1-naphthoxy)propyl]-1-methyl-3-methoxybenzylamine.

5 Compound 22: *N*-(3-phenylpropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 24: *N*-(4-phenylbutyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

10 Compound 34: *N*-(Benzyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 36: *N*-(4-phenoxybutyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 42: *N*-(3-(1-naphthoxy)propyl)-1,1-dimethyl-2-(4-methoxyphenyl).

15 Compound 52: *N*-[2-hydroxy-3-(2-acetamidophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 53: *N*-(2-phenoxyethyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

20 Compound 54: *N*-2-hydroxy-3-(4-*tert*-butylphenoxy)propyl]-1,1-dimethyl-2-phenylethylamine.

Compound 55: *N*-[2-hydroxy-3-(1-naphthoxy)propyl]-1,1-dimethyl-2-phenylethylamine.

Compound 58: *N*-[2-hydroxy-3-(4-acetamidophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

25 Compound 60: *N*-[2-hydroxy-3-(2-phenylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 61: *N*-[2-hydroxy-3-(3-phenylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

30 Compound 62: *N*-[2-Hydroxy-3-(4-phenylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 84: *N*-(2-Phenylethyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

Compound 92: *N*-(2-hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(1-naphthyl)ethylamine.

Compound 93: *N*-(2-hydroxy-3-cyclohexoxypropyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine.

5 Compound 102: *N*-(2-hydroxy-3-phenoxypropyl)-1,1-dimethyl-2-(4-ethoxyphenyl)ethylamine.

Compound 132: *N*-[2-hydroxy-3-(3,4-dimethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, *m/z* (rel. int.) 390 (*M*⁺, .0), 269
10 (17), 268 (100), 163 (6), 153 (5), 121 (21), 114 (17), 77 (5), 71 (19), 70 (17), 58 (7).

Compound 133: *N*-[2-hydroxy-3-(3,5-dimethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, *m/z* (rel. int.) 390 (*M*⁺, .0), 269
15 (16), 268 (100), 193 (9), 163 (8), 154 (7), 121 (24), 114 (46), 76 (6), 71 (11), 70 (18).

Compound 134: *N*-[2-hydroxy-3-(2-carbomethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; ¹H NMR (CDCl₃) δ 1.42 (s, 3H), 1.44 (s, 3H),
20 3.2 (dd, 2H), 3.3-3.6 (bm, 2H), 3.7 (s, 3H), 3.8 (s, 3H), 4.1-4.4 (m, 3H), 4.7 (m, 1H), 6.8 (d, 2H), 7.0 (m, 2H), 7.2 (d, 2H), 7.5 (t, 1H), 7.8 (d, 1H), 8.8 (m, 1H), 9.3 (m, 1H).

Compound 135: *N*-[2-hydroxy-3-(4-methylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
25 GC/EI-MS, *m/z* (rel. int.) 328 (*M*-15, .1), 223 (15), 222 (100), 163 (6), 147 (6), 121 (23), 114 (9).

Compound 136: *N*-[2-hydroxy-3-(2,3-dichlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, *m/z* (rel. int.) 382 (*M*-15, .1), 280
30 (10), 279 (9), 278 (62), 276 (100), 163 (11), 121 (28).

Compound 137: *N*-[2-hydroxy-3-(3,5-dichlorophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, *m/z* (rel. int.) 382 (*M*-15, .1), 280

(11), 279 (9), 278 (65), 277 (16), 276 (100), 163 (8), 146 (5), 144 (5).

Compound 138: N-[2-hydroxy-3-(2,4-dichloro-phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
 5 Hydrochloride; GC/EI-MS, m/z (rel. int.) 382 (M-15,1), 280 (11), 279 (9), 278 (65), 277 (5), 276 (100), 163 (10), 161 (6), 132 (5).

Compound 139: N-[2-hydroxy-3-(3,4-dichloro-phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
 10 Hydrochloride; GC/EI-MS, m/z (rel. int.) 382 (M-15,.1), 279 (10), 279 (9), 278 (63), 276 (100), 163 (9), 146 (5), 121 (29).

Compound 140: N-[2-hydroxy-3-(2,5-dichloro-phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
 15 Hydrochloride; GC/EI-MS, m/z (rel. int.) 382 (M-15,.1), 280 (11), 279 (9), 278 (64), 276 (100), 163 (9), 161 (5), 121 (29), 113 (8).

Compound 141: N-[2-hydroxy-3-(4-ethylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 20 GC/EI-MS, m/z (rel. int.) 342 (M-15,.1), 237 (15), 236 (100), 163 (5), 121 (19), 114 (7).

Compound 142: N-[2-hydroxy-3-(2-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 GC/EI-MS, m/z (rel. int.) 339 (M-15,1), 234 (15), 233 (100),
 25 163 (5), 121 (14).

Compound 143: N-[2-hydroxy-3-(3-nitrophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 GC/EI-MS, m/z (rel. int.) 359 (M-15,.1), 254 (15), 253 (100),
 163 (5), 121 (15).

Compound 144: N-[2-hydroxy-3-(4-ethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 GC/EI-MS, m/z (rel. int.) 358 (M-15,.1), 253 (17), 252 (100),
 30 163 (6), 121 (21), 114 (8), 108 (8).

Compound 145: N-[2-hydroxy-3-(4-iso-propyl-phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.) 356 (M-15, 1), 251 (18), 250 (100), 163 (7), 121 (23), 117 (7), 114 (8).

5 Compound 146: N-[2-hydroxy-3-(3-iso-propyl-phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.) 356 (M-15, 1), 251 (18), 250 (100), 163 (6), 121 (21), 117 (5), 114 (10), 91 (9).

10 Compound 147: N-[2-hydroxy-3-(3-ethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.) 358 (M-15, 1), 253 (16), 252 (100), 163 (5), 121 (20), 114 (12), 77 (5).

Compound 148: N-[2-hydroxy-3-(2-n-propylphenoxy)propyl]-
15 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.) 356 (M-15, 1), 251 (17), 250 (100), 163 (6), 121 (34), 114 (9), 107 (6), 90 (17), 78 (9), 77 (10), 71 (17).

Compound 149: N-[2-hydroxy-3-(4-n-propylphenoxy)propyl]-
20 1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.) 356 (M-15, 8), 251 (18), 250 (100), 163 (5), 121 (19), 114 (7), 110 (5), 107 (7), 91 (6).

Compound 150: N-[2-Hydroxy-3-(3-ethylphenoxy)propyl]-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
25 GC/EI-MS, m/z (rel. int.) 342 (M-15, 6), 237 (16), 236 (100), 163 (5), 121 (21), 114 (6), 105 (5), 90 (6).

Compound 151: N-[2-hydroxy-3-(2-ethylphenoxy)propyl]-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
GC/EI-MS, m/z (rel. int.) 342 (M-15, 1), 237 (16), 236 (100),
30 163 (5), 121 (17), 114 (7), 91 (6), 77 (7).

Compound 152: N-[2-hydroxy-3-(4-trifluoromethoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxy-

phenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.)
398 (M-15,2), 293 (15), 292 (100), 121 (20), 77 (5).

Compound 153: N-[2-hydroxy-3-(2-iso-propyl-
phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
5 Hydrochloride; GC/EI-MS, m/z (rel. int.) 356 (M-15,1), 251
(19), 250 (100), 163 (5), 122 (5), 121 (53), 114 (8), 104
(6), 103 (6), 91 (24), 77 (14).

Compound 154: N-[2-hydroxy-3-(3-trifluoro-
methoxyphenoxy)propyl]-1,1-dimethyl-2-(4-methoxy-
10 phenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.)
398 (M-15,.1), 293 (15), 292 (100), 163 (7), 121 (18).

Compound 155: N-[2-hydroxy-3-(2,6-dichloro-
phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
Hydrochloride; GC/EI-MS, m/z (rel. int.) 382 (M-15,1), 279
15 (11), 279 (9), 277 (64), 275 (100), 163 (11), 163 (5), 161
(6), 121 (33), 114 (12).

Compound 156: N-[2-hydroxy-3-(3,5-bistrifluoro-
methylphenoxy)propyl]-1,1-dimethyl-2-(4-methoxy-
phenyl)ethylamine Hydrochloride; GC/EI-MS, m/z (rel. int.)
20 450 (M-15,.1), 345 (16), 344 (100), 213 (5), 163 (8), 121
(20).

Compound 157: N-[2-hydroxy-3-(3-chloro-5-methoxy-
phenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine
Hydrochloride; GC/EI-MS, m/z (rel. int.) 378 (M-15,.1), 275
25 (5), 274 (34), 273 (16), 272 (100), 163 (5), 121 (17), 114
(8).

Compound 158: N-[2-hydroxy-3-(4-nitrophenoxy)propyl]-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
GC/EI-MS, m/z (rel. int.) 359 (M-15,1), 254 (14), 253 (100),
30 121 (12).

Compound 159: N-[2-hydroxy-3-(2-nitrophenoxy)propyl]-
1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;

GC/EI-MS, m/z (rel. int.) 359 (M-15,.1), 254 (15), 253 (100), 163 (6), 121 (17), 114 (10), 96 (5), 78 (5).

Compound 160: N-[2-hydroxy-3-(3-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 5 GC/EI-MS, m/z (rel. int.) 339 (M-15,.1), 234 (15), 233 (100), 121 (21), 102 (7), 90 (6).

Compound 161: N-[2-hydroxy-3-(4-cyanophenoxy)propyl]-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine Hydrochloride;
 GC/EI-MS, m/z (rel. int.) 339 (M-15,1), 234 (16), 233 (100),
 10 163 (5), 121 (15), 102 (5).

Compound 166: N-[2R-Hydroxy-3-[(2-cyanobenzthien-3-yloxy)propyl]-1,1-dimethyl-2-(3,4-dichlorophenyl)ethylamine. Prepared as a hydrochloride salt, MS (ES) m/e 449 [M+H]⁺.

Compound 167: R-1-[1,1 Dimethyl-2-(4-methoxyphenyl)ethylaminol-3-(2'-carbazoloxo)pran-2-ol. Prepared as a
 15 trifluoroacetate salt, MS (ES) m/e 419.2 [M+H]⁺.

Compound 168: N-[2R-Hydroxy-3-[(2-bromopyridinyloxy)propyl]-1,1-dimethyl-2-(4-methoxy)ethylamine. Prepared as a hydrochloride salt, MS (ES) m/e 411,409 [M+H]⁺.

Compound 169: N-(2-hydroxy-3-(3-N,N-dimethylphenoxy)propyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine, GC/MS
 20 251(100), 176(9), 163(5), 138(11), 137(6),(8), 125(10), 121(23), 114(46), 108(6), 77(6), 76(7),(10), 70(14), 42(8).

Compound 170: N-(2-hydroxy-3-(3-phenylphenoxy)propyl)-1,1-dimethyl-2-(4-methoxyphenyl)ethylamine. GC/MS (mt,
 25 0.1), 284(100), 121(28), 285 (27), 152(13), 70(13), 71(11), 153(10).

Other embodiments are within the following claims.
 Thus, while several embodiments have been shown and
 30 described, various modifications may be made, without departing from the spirit and scope of the present invention.